

RESULT 3

T99173

hypothetical protein T20N10.250 - Arabidopsis thaliana

C:Species: Arabidopsis thaliana (mouse-ear cress)

C:Date: 02-Jun-2000 #sequence_revision 02-Jun-2000 #text_change 08-Dec-2000

C:Accession: T49173

R:ID: Angelo, M.; Vezzi, A.; Modesto, D.; Pigazzi, M.; Valle, G.; Mewes, H.W.; Rudd, S.; I

submitted to the Protein Sequence Database, April 2000

A:Reference number: 225017

A:Accession: T49173

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-517 <DNA>

A:Cross-references: EMBL:AL353032; GSPDB:GN00061; ATSP:T20N10.250

A:Experimental source: cultivar Columbia; BAC clone T20N10

C:Genetics:

A:Gene: ATSP:T20N10.250

A:Map position: 3

A:Introns: 312/3; 359/3; 444/3

C:Superfamily: Arabidopsis thaliana hypothetical protein F17J16.30

Query Match 91.7%; Score 100; DB 2; Length 517;

Best Local Similarity 100.0%; Pred. No. 0.003;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKKKK 21

DB 444 KKKKKKKKKKKKKKKKKKKKK 463

RESULT 4

T18513

hypothetical protein C0845C - malaria parasite (Plasmodium falciparum)

C:Species: Plasmodium falciparum

C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 20-Jun-2000

C:Accession: T18513

R:Lawson, D.; Bowman, S.; Barrett, B.

submitted to the EMBL Data Library, August 1997

A:Reference number: 218935

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-166 <LAW>

A:Cross-references: EMBL:Z98551; PIDN:CAB1123.2

C:Genetics:

A:Map position: 3

A:Introns: 19/1

A>Note: C0845C

Query Match 85.3%; Score 93; DB 2; Length 166;

Best Local Similarity 90.0%; Pred. No. 0.0067;

Matches 18; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKKKK 21

DB 37 KKKKKKKKKKKKKKKKKKKKK 56

RESULT 5

F1619

hypothetical protein PFB0235W - malaria parasite (Plasmodium falciparum)

C:Species: Plasmodium falciparum

C:Date: 13-Nov-1998 #sequence_revision 13-Nov-1998 #text_change 21-Jul-2000

C:Accession: F1619

R:Garner, M.J.; Tetteh, H.; Carnucci, D.J.; Cummings, L.M.; Aravind, L.; Koonin, E.V.;

Peterson, M.; Salberg, S.; Zhou, L.; Sutton, G.G.; Clayton, R.; White, O.; Smith, H.O.

Science 282, 1126-1132, 1998

A:Title: Chromosome 2 sequence of the human malaria parasite Plasmodium falciparum.

A:Reference number: A71600; MUID:99021743; PMID:9804551

A:Accession: F1619

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-483 <GAR>

A:Cross-references: GB:AE001382; GB:AE001362; NID:G3845130; PIDN:AACT1836.1; PID:G3845131

A:Experimental source: clone 3D7

C:Genetics:

A:Gene: PFB0235W

Query Match 82.6%; Score 90; DB 2; Length 483;

Best Local Similarity 85.0%; Pred. No. 0.026;

Matches 17; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKKKK 21

DB 449 KKKKKKKKKKKKKKKKKKKKK 468

RESULT 6

C86477

protein F1504.29 [imported] - Arabidopsis thaliana

C:Species: Arabidopsis thaliana (mouse-ear cress)

C:Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #text_change 31-Mar-2001

C:Accession: C86477

R:Rthellogis, A.; Ecker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, O.; Alonso,

Chen, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Conway, A.R.; Creasy, T.H.; Dewar, K.;

ansen, N.F.; Hughes, B.; Hultzar, L.

Nature 408, 816-820, 2000

A:Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; Kim, C.;

C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, S.X.; Liu, Z.A.; Luros, J.S.; Maiti, R.; Marziani,

Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.

A:Authors: Salberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Tallon, I.

ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.

A:Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.

A:Reference number: A86141; MUID:21016719; PMID:11130712

A:Accession: C86477

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-107 <STD>

A:Cross-references: GB:AE005172; NID:G8778346; PIDN:AAF79354.1; GSPDB:GN00141

C:Genetics:

A:Gene: F1504.29

A:Map position: 1

Query Match 79.8%; Score 87; DB 2; Length 107;

Best Local Similarity 94.4%; Pred. No. 0.019;

Matches 17; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKKKK 19

DB 29 KKKKKKKKKKKKKKKKKKKKK 46

RESULT 7

A48455

acidic phosphoprotein PC5MA1q - Plasmodium chabaudi

C:Species: Plasmodium chabaudi

C:Date: 01-Dec-1993 #sequence_revision 18-Nov-1994 #text_change 20-Mar-1998

C:Accession: A48455

R:Deleersnijder, W.; Prasomitti, P.; Tungpradubkul, S.; Hendrix, D.; Hamers-Casterman, C.

Mol. Biochem. Parasitol. 56, 59-68, 1992

A:Title: Structure of a Plasmodium chabaudi acidic phosphoprotein that is associated with

A:Reference number: A48455; MUID:93116806; PMID:1475002

A:Accession: A48455

A:Status: preliminary

A:Molecule type: nucleic acid

A:Residues: 1-441

A:Cross-references: GB:M95789; NID:G160602; PID:G160603

A:Experimental source: IP-PC1/C

A>Note: sequence extracted from NCBI backbone (NCBIN:121415, NCBI:P:121416)

C:Keywords: phosphoprotein

Query Match 77.1%; Score 84; DB 2; Length 441;

Best Local Similarity 80.0%; Pred. No. 0.089;

Matches 16; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

Oy 2 KKKKKKKKKKKKKKKKKKK 21
| | | | | : | | | | : | | | | :
Db 394 KKKKKKKKKKKKKKKKKKE 413

RESULT 8

hypochemical protein C0425w - malaria parasite (Plasmodium falciparum)
C/Species: Plasmodium falciparum
C/Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 09-Jun-2000
C/Accession: T18440
R/Lawson, D.; Bowman, S.; Barrell, B.
submitted to the EMBL Data Library, August 1997
A/Reference number: Z18935
A/Accession: T18440
A/Status: preliminary; translated from GB/EMBL/DBJ
A/Molecule type: DNA
A/Residues: 1-4550 <LAW>
A/Cross-references: EMBL:Z98547; NID:e1325376; PID:e1325396; PIDN:CA11121.1
C/Genetics:
A/Map position: 3
A/Note: C0425w

Query Match 74.3%; Score 81; DB 2; Length 4550;
Best Local Similarity 44.2%; Pred. No. 0.77;
Matches 19; Conservative 2; Mismatches 0; Indels 22; Gaps 1;

Oy 1 CCKK-----KKKKKKKKKKKKKKKK 21
| | | | | : | | | | | : | | | | | :
Db 132 CCKKNVFNINIKNYFNEKYOTNNIKKKKKKKKKKKKKKK 174

RESULT 9

hypochemical protein C0560c - malaria parasite (Plasmodium falciparum)
C/Species: Plasmodium falciparum
C/Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 09-Jun-2000
C/Accession: T18452
R/Lawson, D.; Bowman, S.; Barrell, B.
submitted to the EMBL Data Library, November 1998
A/Reference number: Z18937
A/Accession: T18452
A/Status: preliminary; translated from GB/EMBL/DBJ
A/Molecule type: DNA
A/Residues: 1-784 <LAW>
A/Cross-references: EMBL:AL008970; NID:e1407852; PID:e1332545; PIDN:CA15594.1
C/Genetics:
A/Map position: 3
A/Note: C0560c

Query Match 73.4%; Score 80; DB 2; Length 784;
Best Local Similarity 75.0%; Pred. No. 0.31;
Matches 15; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

Oy 2 KKKKKKKKKKKKKKKKKKK 21
| | | | | : | | | | | : | | | | | :
Db 360 EKKKKKKKKKKKKKKKKKK 379

RESULT 10

T50609
hypochemical protein DKFZp761B2423.1 - human (fragment)
C/Species: Homo sapiens (man)
C/Date: 21-Jul-2000 #sequence_revision 21-Jul-2000 #text_change 21-Jul-2000
C/Accession: T50609
R/Bloeker, H.; Boecher, M.; Brandt, P.; Mewes, H.W.; Well, B.; Wiemann, S.
submitted to the Protein Sequence Database, June 2000
A/Reference number: Z25143
A/Accession: T50609
A/Status: preliminary
A/Molecule type: mRNA
A/Residues: 1-529 <AAA>
A/Cross-references: EMBL:AL359564

A/Experimental source: adult amygdala; clone DKFZp761B2423
C/Genetics:
A/Note: DKFZp761B2423.1

Query Match 71.6%; Score 78; DB 2; Length 529;
Best Local Similarity 80.0%; Pred. No. 0.37;
Matches 16; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

Oy 2 KKKKKKKKKKKKKKKKKKK 21
| | | | | : | | | | | : | | | | | :
Db 464 KKKKKKKKKKKKKKKKKKK 483

RESULT 11

SAR DNA-binding protein-1 - garden pea
C/Species: Pisum sativum (garden pea)
C/Date: 30-Apr-1999 #sequence_revision 30-Apr-1999 #text_change 02-Jun-2000
C/Accession: T06377
R/Hatton, D.; Gray, J.C.
submitted to the EMBL Data Library, April 1998
A/Description: cDNA encoding a pea SAR DNA-binding protein that shows homology to nucleol
A/Reference number: Z15637
A/Accession: T06377
A/Status: preliminary; translated from GB/EMBL/DBJ
A/Molecule type: mRNA
A/Residues: 1-560 <HAT>
A/Cross-references: EMBL:AF061962; NID:g3132695; PIDN:AA16330.1; PID:g3132696
C/Genetics:
A/Gene: SARBP-1
C/Superfamily: garden pea SAR DNA-binding protein

Query Match 71.6%; Score 78; DB 2; Length 560;
Best Local Similarity 75.0%; Pred. No. 0.38;
Matches 15; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Oy 2 KKKKKKKKKKKKKKKKKKK 21
| | | | | : | | | | | : | | | | | :
Db 463 KKKKKKKKKKKKKKKKKKK 482

RESULT 12

T42727
proliferation potential-related protein - mouse
C/Species: Mus musculus (house mouse)
C/Date: 11-Jan-2000 #sequence_revision 11-Jan-2000 #text_change 02-Sep-2000
C/Accession: T42727
R/Witte, M.M.; Scott, R.B.
submitted to the EMBL Data Library, November 1998
A/Reference number: Z22246
A/Accession: T42727
A/Status: preliminary; translated from GB/EMBL/DBJ
A/Molecule type: mRNA
A/Residues: 1-1560 <WIT>
A/Cross-references: EMBL:U83913; NID:g3858884; PID:g3858885; PIDN:AA172432.1
A/Experimental source: strain Balb/C
C/Genetics:
A/Gene: P2P-R
C/Function:
A/Description: involved in hnRNP association and Rb1 binding
C/Superfamily: RING finger homology
P:57-107/Domain: RING finger homology <RRN>

Query Match 71.6%; Score 78; DB 2; Length 1560;
Best Local Similarity 80.0%; Pred. No. 0.74;
Matches 16; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

Oy 2 KKKKKKKKKKKKKKKKKKK 21
| | | | | : | | | | | : | | | | | :
Db 1497 KKKKKKKKKKKKKKKKKKK 1516

RESULT 13

JC7219

nuclear protein SR-25 - mouse

C:Species: Mus musculus (house mouse)

C>Date: 09-Jun-2000 #sequence_revision 09-Jun-2000 #ext_change 21-Jul-2000

C:Accession: JC7219

R:Saishara, K.; Yamaoka, T.; Moritani, M.; Tanaka, M.; Iwahana, H.; Yoshimoto, K.; Miyag

Biochem. Biophys. Res. Commun. 269, 444-450, 2000

A>Title: Molecular cloning and expression analysis of a putative nuclear protein, SR-25.

A:Reference number: JC7219; MUID:20175222; PMID:10708573

A:Accession: JC7219

A:Molecule type: mRNA

A:Residues: 1-229 <SAS>

A:Cross-references: DDBJ:AB035383; NID:G7619895; PIDN:BA94743.1; PID:G7619896

A:Experimental source: Mink cell line

A:Comment: This protein is a highly hydrophilic nuclear protein with a serine-arginine r

A:Keywords: nucleus; RNA processing

Query Match 70.6%; Score 77; DB 2; Length 229;

Best Local Similarity 75.0%; Pred. No. 0.27;

Matches 15; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKK 21

DB 95 KKKKKKKKKKKKKKKKKKK 114

RESULT 14

A86288

protein F9L1.30 [Imported] - Arabidopsis thaliana

C:Species: Arabidopsis thaliana (mouse-ear cress)

C>Date: 02-Mar-2001 #sequence_revision 02-Mar-2001 #ext_change 31-Mar-2001

C:Accession: A86288

R:Theologis, A.; Becker, J.R.; Palm, C.J.; Federspiel, N.A.; Kaul, S.; White, O.; Alonso,

Chin, C.W.; Chung, M.K.; Conn, L.; Conway, A.B.; Conway, A.R.; Creasy, T.H.; Dewar, K.;

ansen, N.F.; Hughes, B.; Huizar, L.

Nature 408, 816-820, 2000

A:Authors: Hunter, J.L.; Jenkins, J.; Johnson-Hopson, C.; Khan, S.; Khaykin, E.; Kim, C.

C.A.; Li, J.H.; Li, Y.; Lin, X.; Liu, Z.A.; Luos, J.S.; Maiti, R.; Marziani,

Rizzo, M.; Rooney, T.; Rowley, D.; Sakano, H.

A:Authors: Salzberg, S.L.; Schwartz, J.R.; Shinn, P.; Southwick, A.M.; Sun, H.; Tallon,

ker, M.; Wu, D.; Yu, G.; Fraser, C.M.; Venter, J.C.; Davis, R.W.

A>Title: Sequence and analysis of chromosome 1 of the plant Arabidopsis.

A:Reference number: A86141; MUID:21016719; PMID:11130712

A:Accession: A86288

A:Status: preliminary

A:Molecule type: DNA

A:Residues: 1-266 <STO>

A:Cross-references: GB:AE005172; NID:G5103832; PIDN:AAD39662.1; GSPDB:GN00141

A:Genetics:

A:Gene: F9L1.30

A:Map position: 1

Query Match 70.6%; Score 77; DB 2; Length 266;

Best Local Similarity 70.0%; Pred. No. 0.3;

Matches 14; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKK 21

DB 231 KKKRRKKKKKKKKKKKK 250

RESULT 15

T18427

hypothetical protein C0335c - malaria parasite (Plasmodium falciparum)

C:Species: Plasmodium falciparum

C>Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #ext_change 09-Jun-2000

C:Accession: T18427

R:Lawson, D.; Bowman, S.; Barrell, B.

submitted to the EMBL Data Library, August 1997

A:Reference number: T18427

A:Accession: T18427

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-3724 <LAM>

A:Cross-references: EMBL:Z98547; NID:el325376; PID:el325379; PIDN:CAB11104.1

C:Genetics:

A:Introns: 307/1; 1545/2

A>Note: C0335c

Query Match 70.6%; Score 77; DB 2; Length 3724;

Best Local Similarity 75.0%; Pred. No. 1.6;

Matches 15; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 CKKKKKKKKKKKKKKKKKK 20

DB 2201 CEIKKKKKKKKKKKKKKK 2220

Search completed: January 30, 2004, 00:26:20
Job time : 8.97183 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 29, 2004, 23:58:27 ; Search time 4.73239 Seconds
(without alignment)

208.681 Million cell updates/sec

Title: US-09-461-684C-1
Perfect score: 109
Sequence: 1 CXXXXXXXXXXXXXXXXXXXX 21

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues
Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_41:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	84	77.1	441	1	PHPA_PLACH
2	76	69.7	511	1	NOPS_YEAST
3	75	68.8	474	1	CBP5_SCHPO
4	75	68.8	504	1	SIK1_YEAST
5	75	68.8	2404	1	SON_MOUSE
6	75	68.8	2426	1	SON_HUMAN
7	74	67.9	414	1	Y694_METUA
8	74	67.9	474	1	CBP5_KLUUA
9	74	67.9	483	1	CBP5_YEAST
10	74	67.9	726	1	BRD3_HUMAN
11	72	66.1	479	1	CBP5_CANAL
12	72	66.1	678	1	GARP_PLAFR
13	71	65.1	351	1	CG79_HUMAN
14	71	65.1	686	1	CNG1_HUMAN
15	70.5	64.7	1411	1	TCOF_HUMAN
16	70	64.2	233	1	YD08_YEAST
17	70	64.2	599	1	HM21_HUMAN
18	70	64.2	684	1	CNG1_RAT
19	70	64.2	683	1	CNG1_MOUSE
20	69.5	63.8	534	1	NOPS_RAT
21	69.5	63.8	2231	1	SEN1_YEAST
22	69	63.3	142	1	YMH8_YEAST
23	69	63.3	167	1	YK20_YEAST
24	69	63.3	723	1	SSRP_DROME
25	69	63.3	843	1	BLVR_BOVIN
26	68.5	62.8	724	1	Y061_CAEEL
27	68	62.4	523	1	DBP3_YEAST
28	68	62.4	1178	1	MANA_YEAST
29	67	61.5	118	1	Y093_CAEEL
30	67	61.5	690	1	CNG1_BOVIN
31	67	61.5	691	1	CNG1_CANFA
32	67	61.5	1002	1	IF2P_YEAST
33	67	61.5	1220	1	IF2P_HUMAN

34	67	61.5	1362	1	BRD4_HUMAN	O60885 homo sapien
35	67	61.5	2280	1	YCF2_OENHO	O9mf2 oenothera h
36	66	60.6	481	1	CBP5_BEMNI	O43100 emeticella
37	66	60.6	487	1	CBP5_ASFPU	O43102 aspergillus
38	66	60.6	667	1	YK11_SCHPO	O13796 schizosach
39	66	60.6	1153	1	A3D1_HUMAN	O14617 homo sapien
40	66	60.6	1240	1	YVJ1_YEAST	P53935 saccharomyc
41	65.5	60.1	508	1	NO60_DROME	O44081 drosophila
42	65.5	60.1	514	1	DKC1_HUMAN	O60832 homo sapien
43	65	59.6	217	1	KSI_HYDAT	P38978 hydra atten
44	65	59.6	271	1	YGSW_YEAST	P53335 saccharomyc
45	65	59.6	320	1	YD33_YEAST	Q12117 saccharomyc

ALIGNMENTS

RESULT 1	PHPA_PLACH	STANDARD	PRT	441 AA.
AC	002752			
DT	01-JUL-1993 (Rel. 26, Created)			
DT	01-JUL-1993 (Rel. 26, Last sequence update)			
DT	01-JUN-1994 (Rel. 29, Last annotation update)			
DE	Acidic phosphoprotein precursor (50 kDa antigen).			
CN	PCEMAL			
OS	Plasmodium chabaudi.			
OC	Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.			
OX	NCBI_TaxID=5825;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	STRAIN-IP-PCJ.			
RX	MEDLINE=93116806; PubMed=1475002;			
RA	Deleersnyder W., Prasomsilti P., Tungradabkul S., Hendrix D.,			
RA	Hamers-Casterman C., Hamers R.;			
RT	"Structure of a Plasmodium chabaudi acidic phosphoprotein that is			
RT	associated with the host erythrocyte membrane.";			
RL	Mol. Biochem. Parasitol. 56:59-68(1992)			
CC	-1- FUNCTION: DURING INFECTION, THIS PHOSPHOPROTEIN PROBABLY MODULATES			
CC	THE STRUCTURE OF THE RED CELL MEMBRANE TO THE ADVANTAGE OF THE			
CC	PARASITE, ALTHOUGH ITS PRECISE FUNCTION IS NOT KNOWN.			
CC	-1- SUBCELLULAR LOCATION: PERIPHERAL MEMBRANE PROTEIN ON THE			
CC	CYTOPLASMIC FACE OF THE HOST ERYTHROCYTE MEMBRANE.			
CC	-1- MISCELLANEOUS: ASSOCIATED WITH THE HOST RED CELL MEMBRANE			
CC	THROUGHOUT THE ENTIRE ERYTHROCYTIC CYCLE.			
CC	-----			
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration			
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CC	entities requires a license agreement (See http://www.isb-sib.ch/announce/			
CC	or send an email to license@isb-sib.ch).			
CC	-----			
DR	EMBL; M95789; AAA29732.1; -.			
DR	PIR; A48455; A48455.			
KW	Phosphorylation; Signal; Antigen; Membrane; Repeat; Erythrocyte.			
FT	SIGNAL	1	15	OR 24 (POTENTIAL).
FT	CHAIN	16	441	ACIDIC PHOSPHOPROTEIN.
FT	DOMAIN	186	313	16 X 8 AA TANDEN REPEATS.
FT	REPEAT	186	193	1-1.
FT	REPEAT	194	201	1-2.
FT	REPEAT	202	209	1-3.
FT	REPEAT	210	217	1-4.
FT	REPEAT	218	225	1-5.
FT	REPEAT	226	233	1-6.
FT	REPEAT	234	241	1-7.
FT	REPEAT	242	249	1-8.
FT	REPEAT	250	257	1-9.
FT	REPEAT	258	265	1-10.
FT	REPEAT	266	273	1-11.
FT	REPEAT	274	281	1-12.
FT	REPEAT	282	289	1-13.

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FT REPEAT 290 297 1-14.
FT REPEAT 298 305 1-15.
FT REPEAT 306 313 1-16.
FT DOMAIN 353 370 2 X 9 AA TANDEM REPEATS.
FT REPEAT 353 360 2-1.
FT REPEAT 361 368 2-2.
FT DOMAIN 371 417 LYS-RICH (BASIC).
FT CAROHND 21 21 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CAROHND 112 112 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 441 AA, 49708 MW, DB85E83E795EE7E5 CR664;

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Query Match 77.1%; Score 84; DB 1; Length 441;
Best Local Similarity 80.0%; Pred. No. 0.032;
Matches 16; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

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```

Qy 2 KKKKKKKKKKKKKKKKKKK 21
Db 394 KKKKKKKKKKKKKKKKK 413

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RESULT 2
NOP5_YEAST STANDARD; PRT; 511 AA.
ID NOP5_YEAST
AC Q12499;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Nucleolar protein NOP58 (Nucleolar protein NOP5).
GN NOP58 OR NOP5 OR YOR310C OR O6108.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycaceae;
OC NCB1_TaxID=4932;
RX STRAIN=972;
RN SEQUENCE FROM N.A. AND CHARACTERIZATION.
RP STRAIN=W303;
RC MEDLINE=98298165; PubMed=9632712;
RA Wu P., Brockebrough J.S., Metcalfe A.C., Chen S., Arts J.P.;
RL Submitted (Aug-1995) to the EMBL/GenBank/DBJ databases.
RT pre-18S rRNA processing in yeast.
RL J. Biol. Chem. 273:16453-16463(1998).
CC -1- FUNCTION: REQUIRED FOR PRE-18S RNA PROCESSING. MAY BIND
CC MICROTUBULES.
CC -1- SUBUNIT: INTERACTS WITH NOP56 AND NOP1.
CC -1- SUBCELLULAR LOCATION: Nuclear; nucleolar.
CC -1- SIMILARITY: BELONGS TO THE NOP5/NOP56 FAMILY.
CC -----
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CC -----
CC EMBL; X90565; CAAG2165.1; -
CC EMBL; 275217; CAAG9630.1; -
CC EMBL; AF056070; AAC39484.1; -
CC FIR; S58322; S58322.
CC SGD; S0005837; NOP58.
CC DR GO; GO:0005754; C:sma11 nucleolar ribonucleoprotein complex; IPI.
CC DR GO; GO:0003754; F:chaperone activity; NAS.
CC DR GO; GO:001066; F:snRNA binding activity; IIA.
CC DR GO; GO:0030490; P:processing of 20S pre-rRNA; IPI.
CC DR GO; GO:000608; P:snNP protein-nucleus import; NAS.
CC DR InterPro; IPR002687; NOP.
CC DR Pfam; PF01798; NOP; 1.
CC DR Prodom; PD004104; NOP; 1.

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KW Ribosome biogenesis; Nuclear protein; rRNA processing.
FT DOMAIN 441 511 ASP/GLU/LYS-RICH.
SQ SEQUENCE 511 AA, 56956 MW, 8A2889448B2A19E2 CR664;

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Query Match 69.7%; Score 76; DB 1; Length 511;
Best Local Similarity 70.0%; Pred. No. 0.21;
Matches 14; Conservative 6; Mismatches 0; Indels 0; Gaps 0;

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Qy 2 KKKKKKKKKKKKKKKKKKK 21
Db 480 KKKKKKKKKKKKKKKKK 499

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RESULT 3
CBF5_SCHPO STANDARD; PRT; 474 AA.
ID CBF5_SCHPO
AC Q14007;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Centromere/microtubule binding protein cbf5 (centromere-binding factor
DE 5) (Nucleolar protein cbf5).
GN CBF5 OR SPAC29A4.04C.
OS Schizosaccharomyces pombe (Fission yeast).
OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OC Schizosaccharomycaceae;
OC NCB1_TaxID=4896;
RX MEDLINE=21846401; PubMed=11859360;
RN SEQUENCE FROM N.A.
RP STRAIN=972;
RC MEDLINE=21846401; PubMed=11859360;
RA Wood V., Gwilliam R., Rajandream M.A., Lyne M., Lyne R., Stewart A.,
RA Sgourou J., Peat N., Hayles J., Baker S., Basham D., Bowman S.,
RA Brooke K., Brown D., Brown S., Chillingworth T., Churcher C.M.,
RA Collins M., Connor R., Cronin A., Davis P., Feltwell T., Fraser A.,
RA Gentles S., Goble A., Hamlin N., Harris P., Hidalgo J., Hodgson G.,
RA Holtroyd S., Hornsby T., Howarth S., Huckle E.J., Hunt S., Jagsels K.,
RA James K., Jones L., Jones M., Leather S., McDonald S., McLean J.,
RA Mooney P., Moule S., Mungall K., Murphy L., Niblett D., Odell C.,
RA Oliver K., O'Neill S., Pearson D., Quail M.A., Rabinowitch E.,
RA Rutherford K., Rutter S., Saunders D., Seeger K., Sharp S.,
RA Skelton J., Simmonds R., Squares R., Squares S., Stevens K.,
RA Taylor K., Taylor R.G., Tivey A., Walsh S.V., Warren T., Whitehead S.,
RA Woodward J., Volckaert G., Aert R., Robben J., Grymopre B.,
RA Welljens I., Vanstreels E., Rieger M., Schaefer M., Mueller-Auer S.,
RA Gabel C., Fuchs M., Fritze C., Holzer E., Moesli D., Hilbert H.,
RA Borzym K., Langer I., Beck A., Leirach H., Reinhardt R., Pohl T.M.,
RA Eger P., Zimmermann W., Wedler H., Wambutt R., Purnelle B.,
RA Goffeau A., Cadieu E., Dreano S., Gloux S., Lelaure V., Mottier S.,
RA Galibert F., Aves S.J., Xiang Z., Hunt C., Moore K., Hurst S.M.,
RA Lucas M., Rochet M., Galliard C., Tallada V.A., Garzon A., Rhode G.,
RA Daga R.R., Cruzado L., Jimenez J., Sanchez M., del Rey F., Benito J.,
RA Dominguez A., Revuelta J.L., Moreno S., Armstrong J., Forsburg S.L.,
RA Cerutti L., Lowe T., McCombie W.R., Paulsen I., Potashkin J.,
RA Shpakovski G.V., Ussery D., Bartell B.G., Nurse P.;
RT "The genome sequence of Schizosaccharomyces pombe."
RL Nature 415:871-880(2002).
CC -1- FUNCTION: BINDS IN VITRO TO CENTROMERES AND MICROTUBULES. IT IS A
CC CENTROMERE DNA-CBF3-BINDING FACTOR AND IS INVOLVED IN MITOTIC
CC CHROMOSOME SEGREGATION. IT IS ESSENTIAL FOR CELL GROWTH (BY
CC SIMILARITY).
CC -1- SUBCELLULAR LOCATION: Nuclear; nucleolar (by similarity).
CC -1- SIMILARITY: BELONGS TO THE TRUB FAMILY OF PSEUDOURIDINE SYNTHASES.
CC -1- SIMILARITY: Contains 1 PUA domain.
CC -----
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CC EMBL; 297210; CAB10131.1; -
DR PIR; T38485; T38485.
DR Genedb_Spomb; SPAC29A4.04c; -
DR InterPro; IPR004602; CbU5.
DR InterPro; IPR002478; PUA.
DR InterPro; IPR004501; Trub_N.
DR InterPro; IPR004521; Unchar_dom_2.
DR Pfam; PF01472; PUA; 1.
DR Pfam; PF01509; Trub_N; 1.
DR SMART; SM00359; PUA; 1.
DR TIGRFAMs; TIGR00425; CBP5; 1.
DR TIGRFAMs; TIGR00451; unchar_dom_2; 1.
DR PROSITE; PS00890; PUA; 1.
KW Microtubules; Centromere; Repeat; Nuclear protein; DNA-binding.
FT DOMAIN 271 346 PUA.
FT 434 468 7 x 3 AA APPROXIMATE TANDEM REPEATS OF
FT REPEAT 443 445 1.
FT REPEAT 450 452 2.
FT REPEAT 454 456 3.
FT REPEAT 457 459 4.
FT REPEAT 460 462 5.
FT REPEAT 463 465 6.
FT REPEAT 466 468 7.
SQ SEQUENCE 474 AA; 53110 MW; B8C9896C5FAEB71 CRC64;

Query Match 68.8%; Score 75; DB 1; Length 474;
Best Local Similarity 73.7%; Pred. No. 0.24;
Matches 14; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

OY 2 KKKKKKKKKKKKKKKKK 20
Db 454 KKKKKKKKKKKKKKKK 472

RESULT 4
SIK1_YEAST STANDARD; PRT; 504 AA.
ID SIK1_YEAST
AC Q12460;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE SIK1 protein (Nucleolar protein NOP56).
GN SIK1 OR NOP56 OR YLR197W OR I6167.9.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=5288C / YPH1;
RX MEDLINE=96040178; PubMed=7547500;
RA Morin P.J., Downs J.A., Snodgrass A.M., Gilmore T.D.;
RT "Genetic analysis of growth inhibition by GAL4-L kappa B-alpha in
RT Saccharomyces cerevisiae."
RL Cell Growth Differ. 6:789-798(1995).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=5288C / AB972;
RX MEDLINE=97313267; PubMed=9169871;
RA Johnson M., Hillier L., Riles L., Albermann K., Andre B., Anseorge W.,
RA Benes V., Brueckner M., Delius H., Dubois E., Duesterhoeft A.,
RA Enlian K.-D., Floeth M., Goffeau A., Hedling U., Heumann K.,
RA Heuss-Neitzel D., Hilbert H., Hilger F., Kleine K., Koelter P.,
RA Louis E.-J., Messenguy F., Mewes H.-W., Miosga T., Moestl D.,
RA Mueller-Auer S., Nettlich U., Obermaier B., Piravandi E., Pohl T.M.,
RA Poterelle D., Purnelle B., Rechmann S., Rieger M., Rinke M., Rose M.,
RA Scharfe M., Scheu B., Scholler P., Schwager C., Schwarz S.,
RA Underwood A.P., Urrestalazu L.A., Vandenbol M., Verhasselt P.,
RA Viereckels F., Voet M., Volckaert G., Voss H., Wambutt R., Wedler E.,
RA Wedler H., Zimmermann F.K., Zollner A., Hant J., Hobeisel J.D.;
RT "The nucleotide sequence of Saccharomyces cerevisiae chromosome XII."

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RL Nature 387:87-90(1997).
RN [3]
RP CHARACTERIZATION, AND MUTAGENESIS.
RX MEDLINE=98038777; PubMed=9372940;
RA Gauthier T., Berges T., Tollervy D., Hurt E.;
RT "Nucleolar KEE/D repeat proteins Nop56p and Nop58p interact with Nop1p
RT and are required for ribosome biogenesis."
RL Mol. Cell. Biol. 17:7088-7098(1997).
CC -1- FUNCTION: REQUIRED FOR 60S RIBOSOMAL SUBUNIT SYNTHESIS.
CC -1- SUBUNIT: INTERACTS WITH NOP1 AND NOP58.
CC -1- SUBCELLULAR LOCATION: Nuclear; nucleolar.
CC -1- SIMILARITY: BELONGS TO THE NOP5/NOP56 FAMILY.
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CC
DR EMBL; U20237; AAC49066.1; -.
DR EMBL; U14913; AAB67431.1; -.
DR PIR; S48550; S48550.
DR SGD; S0004187; SIK1.
DR GO; GO:0005732; C:small nucleolar ribonucleoprotein complex; IPI.
DR GO; GO:0030490; P:processing of 20S pre-rRNA; IPI.
DR InterPro; IPR002687; NOP.
DR Pfam; PF01798; NOP; 1.
DR ProDom; PD004104; NOP; 1.
KW Ribosome biogenesis; Nuclear protein.
FT DOMAIN 443 504
FT MUTAGEN 333 333 ASP/GLU/LYS-RICH.
FT MUTAGEN 333 333 V->A: REDUCED GROWTH RATE AT ALL
FT MUTAGEN 355 355 TEMPERATURES; WHEN ASSOCIATED WITH R-385.
FT MUTAGEN 355 355 Y->C: AT 37 DEGREES, GROWTH SLOWS AFTER 6
FT 20 HOURS AND CELL DIVISION STOPS AFTER
FT 20 HOURS.
FT MUTAGEN 385 385 M->R: REDUCED GROWTH RATE AT ALL
FT SEQUENCE 504 AA; 56864 MW; F8522A5870EF4842 CRC64;

Query Match 68.8%; Score 75; DB 1; Length 504;
Best Local Similarity 70.0%; Pred. No. 0.25;
Matches 14; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

OY 2 KKKKKKKKKKKKKKKKK 21
Db 465 KKKKKKKKKKKKKKKK 464

RESULT 5
SON_MOUSE STANDARD; PRT; 2404 AA.
ID SON_MOUSE
AC Q9QX47; Q9CQ12; Q9COK6; Q9QXPS;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE SON protein.
GN SON.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A. (ISOFORMS 1 AND 2).
RC STRAIN=129/Sv;
RX MEDLINE=2040886; PubMed=10950926;
RA Wynn S.L., Fisher R.A., Pregel C., Price M., Liu Q.Y., Khan I.M.,
RA Zammit P., Dadgar K., Mazzari W., Kessling A., Lee J.S., Bulwella L.;
RT "Organization and conservation of the GART/SON/DONSON locus in mouse
RT and human genomes."
RL Genomics 68:57-62(2000).
RN [2]

```

RP SEQUENC OF 1-116 FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Hippocampus, Small intestine, and Tongue;
RX MEDLINE=21085660; PubMed=11217851;
RA Kawai J., Shinagawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
RA Aizawa K., Izawa M., Niehi K., Kiyosawa H., Kondo S., Yamanka I.,
RA Saito T., Okazaki Y., Gojibori T., Bono H., Kanakawa T., Saito R.,
RA Kadota K., Matsuda H.A., Ashburner M., Batilov S., Caavaant T.,
RA Fletschmann W., Gaeseleland T., Gissi C., King B., Kochiwa H.,
RA Kueth P., Lewis S., Matsumoto Y., Nikiado I., Pesole G., Quackenbush J.,
RA Schiml L.M., Staubli F., Suzuki R., Tomita M., Wegner L., Washio T.,
RA Sakai K., Okido T., Furuno M., Anono H., Baldarelli R., Barsh G.,
RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bulc C., Fletcher C., Fujita M., Gariboldi M.,
RA Gustenich S., Hill D., Hofmann M., Hume D.A., Kanlaya M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima Y., Mazzarelli U., Momberts P.,
RA Norone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
RA Sasaki H., Sato K., Schoenbach C., Seta T., Shibata Y., Storch K.-F.,
RA Suzuki H., Toyooka K., Wang K.H., Weltz C., Whitaker C., Wilmink L.,
RA Wynshaw-Boris A., Yoshida K., Haesgawa Y., Kawai H., Kontsuki S.,
RA Hayashizaki Y.;
RT "Functional annotation of a full-length mouse cDNA collection";
RL Nature 409:685-690(2001).
CC -1- FUNCTION: Transcriptional repressor. Binds to the consensus DNA
CC sequence: 5'-GAC[CT]AN[CG][AG]CC-3'. Might protect cells from
CC apoptosis. Might be involved in pre-mRNA splicing (By similarity).
CC -1- SUBCELLULAR LOCATION: Nuclear (By similarity).
CC -1- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=2;
CC Name=1;
CC IsoId=G9QX47-1; Sequence=Displayed;
CC Name=2;
CC IsoId=G9QX47-2; Sequence=VSD_004416, VSP_004417;
CC -1- TISSUE SPECIFICITY: Widely expressed.
CC -1- DOMAIN: Contains 8 types of repeats which are distributed in 3
CC regions.
CC -1- SIMILARITY: Contains 1 DBM (double-stranded RNA-binding) domain.
CC -1- SIMILARITY: Contains 1 DBM (double-stranded RNA-binding) domain.
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CC -----
CC EMBL: AF193606; AAF23120.1; -;
DR EMBL: AF193595; AAF23120.1; JOINED.
DR EMBL: AF193596; AAF23120.1; JOINED.
DR EMBL: AF193597; AAF23120.1; JOINED.
DR EMBL: AF193598; AAF23120.1; JOINED.
DR EMBL: AF193599; AAF23120.1; JOINED.
DR EMBL: AF193600; AAF23120.1; JOINED.
DR EMBL: AF193601; AAF23120.1; JOINED.
DR EMBL: AF193602; AAF23120.1; JOINED.
DR EMBL: AF193603; AAF23120.1; JOINED.
DR EMBL: AF193604; AAF23120.1; JOINED.
DR EMBL: AF193605; AAF23120.1; JOINED.
DR EMBL: AF193607; AAF23120.1; -;
DR EMBL: AK019312; BAB31659.1; -;
DR EMBL: AK019081; BAB31536.1; -;
DR EMBL: AK008478; BAB25691.1; -;
DR EMBL: AK008256; BAB2562.1; -;
DR MGD: MGI:98353; Son.
DR GO: GO:0005515; P;protein binding activity; IPT.
DR InterPro: IPR001159; DS_RBD.
DR InterPro: IPR000467; G_patch.
DR Pfam: PF00035; dsrm; 1.
DR Pfam: PF01585; G_patch; 1.
DR SMART: SM00443; G_patch; 1.
DR PROSITE: PS50137; DS_RBD; 1.
DR PROSITE: PS50174; G_patch; 1.

KW	RNA-binding; DNA-binding; Nuclear protein; Repeat;
FM	Alternative splicing.
FT	DOMAIN 721 850
FT	DOMAIN 867 943
FT	DOMAIN 961 1080
FT	REPEAT 961 966
FT	REPEAT 969 974
FT	REPEAT 976 981
FT	REPEAT 985 990
FT	REPEAT 993 998
FT	REPEAT 1001 1006
FT	REPEAT 1010 1015
FT	REPEAT 1018 1023
FT	REPEAT 1026 1031
FT	REPEAT 1035 1040
FT	REPEAT 1044 1049
FT	REPEAT 1055 1060
FT	REPEAT 1066 1071
FT	REPEAT 1075 1080
FT	DOMAIN 1101 1133
FT	DOMAIN 1910 1979
FT	REPEAT 1910 1916
FT	REPEAT 1938 1944
FT	REPEAT 1945 1951
FT	REPEAT 1952 1958
FT	REPEAT 1959 1965
FT	REPEAT 1966 1972
FT	REPEAT 1973 1979
FT	DOMAIN 1919 1990
FT	REPEAT 1919 1937
FT	REPEAT 1980 1990
FT	DOMAIN 1991 2017
FT	DOMAIN 2283 2329
FT	DOMAIN 2349 2404
FT	VARSPLIC 2086 2086
FT	VARSPLIC 2087 2404
SQ	SEQUENCE 2404 AA; 261428 MM; 648BF78ED3FC01D9 CRC64;
Query Match	68.8%; Score 75; DB 1; Length 2404;
Best Local Similarity	75.0%; Pred. No. 0.85;
Matches	15; Conservative 2; Mismatches 3; Indels 0; Gaps 0;
OY	2 KKKKKKKKKKKKKKKK 21
DB	109 KKHKKHKKKKKKKKKK 128
RESULT 6	
SON_HUMAN	STANDARD; PRT; 2426 AA.
ID	P15583; Q14487; Q95981; Q14120; Q9H7B1; Q9P070; Q9UKP9;
AC	Q9UPY0;
DT	01-NOV-1990 (Rel. 16, Created)
DT	28-FEB-2003 (Rel. 41, Last sequence update)
DT	28-FEB-2003 (Rel. 41, Last annotation update)
DE	SON protein (SON3) (Negative regulatory element-binding protein) (NRE-binding protein) (DBP-5) (Box antagonist selected in saccharomyces 1) (BASIS) (Protein C2orf50).
DE	SON OR NREBP OR DBP5 OR C2IORF50 OR KIAA1019.
OS	Homo sapiens (Human).
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC	Mammalia; Metascia; Primates; Catarrhini; Hominoidea; Homo.
OX	NCBI_TaxId=9606;
NP	[1]
RP	SEQUENCE FROM N.A. (ISOFORMS A; B; C; D; E AND F).
RX	MEDLINE=21564202; PubMed=11707072;

RA Raymond A., Friedli M., Neergaard Heinrichsen C., Chapot F.,
 RA Deutsch S., Ucla C., Rossier C., Lyle R., Guipponi M.,
 RA Antonarakis S.E.;
 RT "From PREDS and open reading frames to cDNA isolation: revisiting the
 RT human chromosome 21 transcription map."; *Genomics* 78:46-54(2001).
 RL [12]
 RN SEQUENCE FROM N.A. (ISOFORM G).
 RP TISSUE=Liver;
 RC MEDLINE=21316479; PubMed=11306577;
 RX Sun C.-T., Lo W.-Y., Wang I.-H., Lo Y.-H., Shiou S.-R., Lai C.-K.,
 RA Ting L.-P.;
 RT "Transcription repression of human hepatitis B virus genes by negative
 RT regulatory element-binding protein/SOX."; *J. Biol. Chem.* 276:24059-24067(2001).
 RL [3]
 RN SEQUENCE OF 1-689 FROM N.A. (ISOFORM H).
 RP TISSUE=Placenta;
 RC Casadei R., Sciripoli P., D'Addabbo P., Canaider S., Lenzi L.,
 RA Vitale L., Giannone S., Carinci P., Zannotti M.;
 RL Submitted (Oct-2001) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE OF 1-130 FROM N.A.
 RC TISSUE=Smooth muscle;
 RA Kawakami T., Noguchi S., Itoh T., Shigeta K., Senba T., Matsumura K.,
 RA Nakajima Y., Mizuno T., Morinaga M., Tanigami A., Fujiwara T., Ono T.,
 RA Yamada K., Fujii Y., Ozaki K., Hirao M., Ohmori Y., Oca T., Suzuki Y.,
 RA Ohsayashi M., Nishi T., Shibahara T., Tanaka T., Nakamura Y.,
 RA Isogai T., Sugano S.;
 RT "NEBO human cDNA sequencing project."; *Submitted (Aug-2000) to the EMBL/GenBank/DBJ databases.*
 RL [5]
 RN SEQUENCE OF 1-114 FROM N.A.
 RP TISSUE=Blood;
 RC Ye M., Zhang Q.H., Zhou J., Shen Y., Wu X.Y., Guan Z.Q., Wang L.,
 RA Fan H.Y., Mao Y.F., Dai M., Huang Q.H., Chen S.J., Chen Z.;
 RT "Human partial CDS from cd34+ stem cells."; *Submitted (May-1999) to the EMBL/GenBank/DBJ databases.*
 RL [6]
 RN SEQUENCE OF 437-2426 FROM N.A. (ISOFORM B).
 RP TISSUE=Brain;
 RC MEDLINE=93937452; PubMed=10470851;
 RA Kikuno R., Nagase T., Ishikawa K.-I., Hirosewa M., Miyajima N.,
 RA Tanaka A., Kozani H., Nomura N., Ohara O.;
 RT "Prediction of the coding sequences of unidentified human genes. XIV.
 RT The complete sequences of 100 new cDNA clones from brain which code
 RT for large proteins in vitro."; *DNA Res.* 6:197-205(1999).
 RL [7]
 RN SEQUENCE OF 554-2426 FROM N.A. (ISOFORM A).
 RP MEDLINE=92049296; PubMed=1944255;
 RA Chumakov I.M., Berdichevskii F.B., Sokolova N.V., Reznikov M.V.,
 RA Prasolov V.S.;
 RT "Identification of a protein product of a novel human gene SON and
 RT the biological effect upon administering a changed form of this gene
 RT into mammalian cells."; *Mol. Biol.* 25:731-740(1991).
 RL [8]
 RN SEQUENCE OF 709-1079 FROM N.A. (ISOFORM I).
 RP TISSUE=Placenta;
 RC MEDLINE=93062885; PubMed=1435774;
 RX Bliskovskii V.V., Kirillov A.V., Zakhariev V.M., Chumakov I.M.;
 RT "The human son gene: the large and small transcripts contains various
 RT 5'-terminal sequences."; *Mol. Biol.* 26:807-812(1992).
 RL [9]
 RN SEQUENCE OF 1009-1131 FROM N.A.
 RP TISSUE=Placenta;
 RC MEDLINE=93062884; PubMed=1435773;
 RX Bliskovskii V.V., Berdichevskii F.B., Tkachenko A.V., Belova M.E.,
 RA Chumakov I.M.;
 RT "Coding part of the son gene small transcript contains four areas of
 RT complete tandem repeats.";

RL Mol. Biol. (Mosk) 26:793-806(1992).
 RN [10]
 RP SEQUENCE OF 1145-2426 FROM N.A. (ISOFORM F).
 RX MEDLINE=93048367; PubMed=1424386;
 RA Mattioni T., Hume C.R., Konigorski S., Hayes P., Osterweil Z.,
 RA Lee J.S.;
 RT "A cDNA clone for a novel nuclear protein with DNA binding
 RT activity."; *Chromosome* 101:618-624(1992).
 RL [11]
 RN SEQUENCE OF 1692-2175 FROM N.A. (ISOFORM A).
 RP MEDLINE=89039788; PubMed=3054499;
 RX Berdichevskii F.B., Chumakov I.M., Kiselev L.L.;
 RT "Decoding of the primary structure of the son3 region in human
 RT genome: identification of a new protein with unusual structure and
 RT homology with DNA-binding proteins."; *Mol. Biol.* (Mosk) 22:794-801(1988).
 RL [12]
 RN SEQUENCE OF 1939-2426 FROM N.A. (ISOFORM J).
 RP TISSUE=Cerebellum;
 RC MEDLINE=99439804; PubMed=10509013;
 RX Greenhalf W., Lee J., Chaudhuri B.;
 RT "A selection system for human apoptosis inhibitors using yeast.";
 RL Yeast 15:1307-1321(1999).
 CC -1- FUNCTION: Represses hepatitis B virus (HBV) core promoter activity
 CC and transcription of HBV genes and production of HBV virions.
 CC Binds to the consensus DNA sequence: 5'-GA(GT)AN(GG)AG(C)C-3'.
 CC Might protect cells from apoptosis. Might be involved in pre-mRNA
 CC splicing (By similarity).
 CC -1- SUBCELLULAR LOCATION: Nuclear with a speckled distribution.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event-Alternative splicing: Named isoforms=10;
 CC Comment=Experimental confirmation may be lacking for some
 CC isoforms;
 CC Name=F;
 CC IsoId=P18583-1; Sequence=Displayed;
 CC Name=A;
 CC IsoId=P18583-2; Sequence=VSP_004401, VSP_004402, VSP_004403;
 CC Name=B;
 CC IsoId=P18583-3; Sequence=VSP_004404, VSP_004405;
 CC Name=C;
 CC IsoId=P18583-4; Sequence=VSP_004406, VSP_004407;
 CC Name=D;
 CC IsoId=P18583-5; Sequence=VSP_004403;
 CC Name=E;
 CC IsoId=P18583-6; Sequence=VSP_004408, VSP_004409;
 CC Name=G;
 CC IsoId=P18583-7; Sequence=VSP_004410;
 CC Name=H;
 CC IsoId=P18583-8; Sequence=VSP_004411, VSP_004412;
 CC Name=I;
 CC IsoId=P18583-9; Sequence=VSP_004413;
 CC Name=J;
 CC IsoId=P18583-10; Sequence=VSP_004414, VSP_004415;
 CC -1- TISSUE SPECIFICITY: Widely expressed, with the higher expression
 CC seen in leukocyte and heart.
 CC -1- DOMAIN: Contains 8 types of repeats which are distributed in 3
 CC regions.
 CC -1- MISCELLANEOUS: Colocalizes with the pre-mRNA splicing factor
 CC SFRS2/SC-35.
 CC -1- SIMILARITY: Contains 1 G-patch domain.
 CC -1- SIMILARITY: Contains 1 DBM (double-stranded RNA-binding) domain.
 CC -1- CAUTION: ISOFORM A SEQUENCE FROM REF.7 DIFFERS FROM THAT SHOWN DUE
 CC TO A FRAMESHIFT.
 CC -1- CAUTION: ISOFORM F SEQUENCE FROM REF.10 DIFFERS FROM THAT SHOWN
 CC DUE TO A FRAMESHIFT.
 CC -----
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CC EMBL; AF380179; AAL34497.1; -
 DR EMBL; X63753; CAA45282.1; ALT_FRAME.
 DR EMBL; M36438; AAA36624.1; -
 DR EMBL; AF380180; AAL34498.1; -
 DR EMBL; AF380181; AAL34499.1; -
 DR EMBL; AF380182; AAL34500.1; -
 DR EMBL; AF380183; AAL34501.1; -
 DR EMBL; AF380184; AAL34502.1; -
 DR EMBL; AY025895; AAK07692.1; -
 DR EMBL; AF435977; AAL30810.1; -
 DR EMBL; X63751; CAC69885.1; -
 DR EMBL; AB028942; BAA82971.1; -
 DR EMBL; X63071; CAA44793.1; ALT_FRAME.
 DR EMBL; AF139897; AAD50078.1; -
 DR EMBL; AK024752; BAB14985.1; -
 DR EMBL; AF161428; AAF28988.1; -
 DR EMBL; AF161430; AAF28990.1; -
 DR Genew; HGNC:11183; SON.
 DR GK; P18583; -
 DR MIM; 182465; -
 DR GO; GO:0008189; F.apoptosis inhibitor activity; IDA.
 DR GO; GO:0003677; F.DNA binding activity; TAS.
 DR GO; GO:0006916; P.apoptosis; IDA.
 DR InterPro; IPR001159; DS_RBD.
 DR InterPro; IPR000467; G_patch.
 DR Pfam; PF00035; dsrm; 1.
 DR Pfam; PF01585; G_patch; 1.
 DR SMART; SM00358; DSRM; 1.
 DR SMART; SM00443; G_patch; 1.
 DR PROSITE; PS0137; DS_RBD; 1.
 DR PROSITE; PS0174; G_PATCH; 1.
 KW RNA-binding; DNA-binding; Nuclear protein; Repeat;
 KW Alternative splicing.
 FT DOMAIN 726 895 17 X 10 AA TANDEM REPEATS OF L-A-[ST]-
 [NSG]-[TS]-MDSQM.

Query Match 68.8%; Score 75; DB 1; Length 2426;
 Best Local Similarity 75.0%; Pred. No. 0.86;
 Matches 15; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

Qy 2 KKKKKKKKKKKKKKKKKKK 21
 Db 109 KKKKKKKKKKKKKKKKKKK 128

RESULT 7
 Y694 METUA STANDARD; PRT; 414 AA.
 ID Y694 METUA
 AC 058105;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Hypothetical protein Mj0694.
 GN Mj0694.
 OS Methanococcus jannaschii.
 OC Archaea; Euryarchaeota; Methanococci; Methanococcales;
 OC Methanocaldococcaceae; Methanocaldococcus.
 OX NCBI_TaxId=2190;
 RN (1)
 RP SEQUENCE FROM N.A.
 RC STRAIN=JAL-1 / DSM 2661 / ATCC 43067;
 RX MEDLINE=96337999; PubMed=868087;
 RA Bult C.V., White O., Olsen G.J., Zhou L., Fleischmann R.D.,
 RA Sutton G.G., Blake J.A., Fitzgerald L.M., Clayton R.A., Gocayne J.D.,
 RA Kerlavage A.R., Dougherty B.A., Tomb J.-F., Adams M.D., Reich C.I.,
 RA Overbeek R., Kirkness E.F., Weissbrock K.G., Merrick J.M., Glodek A.,
 RA Scott J.L., Geoghegan N.S.M., Weidman J.F., Fuhmann J.L., Nguyen D.,
 RA Cutlerback T.R., Kelley J.M., Peterson J.D., Sadow P.W., Hanna M.C.,
 RA Cotton M.D., Roberts K.M., Hurst M.A., Kaine B.P., Borodovsky M.,
 RA Klenk H.-P., Fraser C.M., Smith H.O., Woese C.R., Venter J.C.;
 RA "Complete genome sequence of the methanogenic archaeon, Methanococcus

RT jannaschii.";
 RL Science 273:1058-1073 (1996).
 CC -1- SIMILARITY: BELONGS TO THE NOP5/NOP56 FAMILY.
 CC
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CC EMBL; U67516; AAB98689.1; -
 DR PIR; F64386; F64386.
 DR TIGR; Mj0694; -
 DR InterPro; IPR002687; NOP.
 DR Pfam; PF01798; NOP; 1.
 DR Prodom; PD004104; NOP; 1.
 DR Hypothetical protein; Complete proteome.
 KW Domain 349 414 ASP/GLU/LYS-RICH.
 FT SEQUENCE 414 AA; 47799 MW; A9092EFC3C82C407 CRC64;

Query Match 67.9%; Score 74; DB 1; Length 414;
 Best Local Similarity 75.0%; Pred. No. 0.27;
 Matches 15; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

Qy 2 KKKKKKKKKKKKKKKKKKK 21
 Db 377 KKKKKKKKKKKKKKKKKKK 396

RESULT 8
 CBFS_KLULA STANDARD; PRT; 474 AA.
 ID CBFS_KLULA
 AC 013473;
 DT 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Centromere/microtubule binding protein CBFS (Centromere-binding factor
 DE 5) (Nucleolar protein CBFS).
 GN CBFS.
 OS Kluveromyces lactis (Yeast).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 OC Saccharomycetales; Saccharomycetaceae; Kluveromyces.
 OX NCBI_TaxId=26985;
 RN (1)
 RP SEQUENCE FROM N.A.
 RC STRAIN=JBD100;
 RX MEDLINE=96144788; PubMed=9483794;
 RA Winkler A.A., Bobok A., Zonneveld B.J.M., Steensma H.Y.,
 RA Hooykaas P.J.J.;
 RT "The lysine-rich C-terminal repeats of the centromere-binding factor
 RT 5 (Cbfs) of Kluveromyces lactis are not essential for function.";
 RL Yeast 14:37-48 (1998).
 CC -1- FUNCTION: BINDS IN VITRO TO CENTROMERES AND MICROTUBULES. IT IS A
 CC CENTROMERE DNA-CBFS-BINDING FACTOR AND IS ESSENTIAL FOR CELL GROWTH. MAY BE IN
 CC CHROMOSOME SEGREGATION. IT IS ESSENTIAL FOR CELL GROWTH. MAY BE IN
 CC SOME WAY ASSOCIATED WITH THE CBFS 110 kDa SUBUNIT (CBFS3) (BY
 CC SIMILARITY).
 CC -1- SUBCELLULAR LOCATION: Nuclear; nucleolar (By similarity).
 CC -1- SIMILARITY: BELONGS TO THE TRUB FAMILY OF PSEUDOURIDINE SYNTHASES.
 CC -1- SIMILARITY: Contains 1 PUA domain.
 CC
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CC EMBL; AF008563; AAC64862.1; -
 DR InterPro; IPR004802; CBFS.

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DR InterPro; IPR002478; PUA.
DR InterPro; IPR002501; Trub_N.
DR InterPro; IPR004521; Unchar_dom_2.
DR Pfam; PF01472; PUA; 1.
DR Pfam; PF01509; Trub_N; 1.
DR SMART; SM00359; PUA; 1.
DR TIGRFAMs; TIGR00425; CBFS; 1.
DR TIGRFAMs; TIGR00451; unchar_dom_2; 1.
DR PROSITE; PS00890; PUA; 1.
KM Microtubules; Centromere; Repeat; Nuclear protein; DNA-binding.
FT DOMAIN 265 340 PUA.
FT DOMAIN 431 460 1.
FT REPEAT 431 433 1.
FT REPEAT 434 436 2.
FT REPEAT 437 439 3.
FT REPEAT 440 442 4.
FT REPEAT 443 445 5.
FT REPEAT 446 448 6.
FT REPEAT 449 451 7.
FT REPEAT 452 454 8.
FT REPEAT 455 457 9.
SQ SEQUENCE 474 AA; 53630 MW; 95306EC7E7EA756C CRC64;

Query Match 67.9%; Score 74; DB 1; Length 474;
Best Local Similarity 70.0%; Pred. No. 0.3;
Matches 14; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

OY 2 KKKKKKKKKKKKKKKKKKKKK 21
DB 434 KKKKKKKKKKKKKKKKKKK 453

RESULT 9
CBFS_YEAST STANDARD; PRT; 483 AA.
ID CBFS_YEAST
AC P33322;
FT 01-FEB-1994 (Rel. 28, Created)
DT 01-FEB-1994 (Rel. 28, Last sequence update)
DE 15-SEP-2003 (Rel. 42, Last annotation update)
DE Centromere/microtubule binding protein CBFS (Centromere-binding factor 5) (Nucleolar protein CBFS) (P64').
GN CBFS OR YLR175W OR U9470.11.
OS Saccharomyces cerevisiae (Baker's Yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
OC NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.
RX MEDLINE=93330283; PubMed=8336724;
RA Jiang W., Middleton K., Yoon H.-J., Fouquet C., Carbon J.;
RA "An essential yeast protein, CBFSp, binds in vitro to centromeres and microtubules."
RT Mol. Cell. Biol. 13:4884-4893(1993).
RL [2]
RM SEQUENCE FROM N.A.
RC STRAIN=S288c / AB972;
RX MEDLINE=93713267; PubMed=9169871;
RA Johnston M., Hillier L., Riles L., Albermann K., Andre B., Ansgore W.,
RA Benes V., Brueckner M., Delius H., Dubois E., Duesterhoeft A.,
RA Eutana K.-D., Floeth M., Goffeau A., Hebling U., Heumann K.,
RA Heuss-Neitzel D., Hilbert H., Hilger F., Kline K., Koetter P.,
RA Louis E.J., Messenguy F., Mewes H.-W., Miosga T., Moestl D.,
RA Mueller-Auer S., Netewich U., Obermayer B., Piravandi E., Pohl T.M.,
RA Portellelle D., Purnelle B., Recheimann S., Rieger M., Rinke M., Rose M.,
RA Scharfe M., Scherens B., Scholler P., Schwaiger C., Schwarz S.,
RA Underwood A.P., Urrutazu L.A., Vandenbol M., Verhaesselt P.,
RA Viereckels F., Voet M., Volckaert G., Voss H., Wambutt R., Wedler E.,
RA Wedler H., Zimmermann F.K., Zollner A., Hani J., Hobeisel J.D.,
RA "The nucleotide sequence of Saccharomyces cerevisiae chromosome XII."
RT Nature 387:87-90(1997).
RL -1- FUNCTION: BINDS IN VITRO TO CENTROMERES AND MICROTUBULES. IT IS A
CC CENTROMERE DNA-CBF3-BINDING FACTOR AND IS INVOLVED IN MITOTIC
CC CHROMOSOME SEGREGATION. IT IS ESSENTIAL FOR CELL GROWTH. MAY BE IN

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CC SOME WAY ASSOCIATED WITH THE CBF3 110 KDa SUBUNIT (CBF3A).
CC -1- SUBCELLULAR LOCATION: Nuclear; nucleolar.
CC -1- SIMILARITY: BELONGS TO THE TRUB FAMILY OF PSEUDOURIDINE SYNTHASES.
CC -1- SIMILARITY: Contains 1 PUA domain.
CC -----
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CC -----
DR EMBL; L12351; AAA34473.1; -
DR EMBL; U17246; AAB67463.1; -.
DR PIR; S41853; S41853.
DR SGD; S0004165; CBFS.
DR GO; GO:0005732; C:small nucleolar ribonucleoprotein complex; IPI.
DR InterPro; IPR004802; Cbf5.
DR InterPro; IPR002478; PUA.
DR InterPro; IPR002501; Trub_N.
DR InterPro; IPR004521; Unchar_dom_2.
DR Pfam; PF01472; PUA; 1.
DR Pfam; PF01509; Trub_N; 1.
DR SMART; SM00359; PUA; 1.
DR TIGRFAMs; TIGR00451; unchar_dom_2; 1.
DR PROSITE; PS00890; PUA; 1.
KM Microtubules; Centromere; Repeat; Nuclear protein; DNA-binding.
FT DOMAIN 266 341 PUA.
FT DOMAIN 434 463 1.
FT REPEAT 434 436 10 X 3 AA TANDEM REPEATS OF K-K [DE].
FT REPEAT 437 439 1.
FT REPEAT 440 442 2.
FT REPEAT 443 445 3.
FT REPEAT 446 448 4.
FT REPEAT 449 451 5.
FT REPEAT 452 454 6.
FT REPEAT 455 457 7.
FT REPEAT 458 460 8.
FT REPEAT 461 463 9.
SQ SEQUENCE 483 AA; 54704 MW; D356B39FDC32E2D CRC64;

Query Match 67.9%; Score 74; DB 1; Length 483;
Best Local Similarity 70.0%; Pred. No. 0.3;
Matches 14; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

OY 2 KKKKKKKKKKKKKKKKKKKKK 21
DB 434 KKKKKKKKKKKKKKKKKKK 453

RESULT 10
BRD3_HUMAN STANDARD; PRT; 726 AA.
ID BRD3_HUMAN
AC Q15059; Q92645;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DE 16-OCT-2001 (Rel. 40, Last annotation update)
DE Bromodomain-containing protein 3 (RING3-like protein).
GN BRD3 OR RING3L OR KIAA0043.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OC NCBI_TaxID=9606;
RN [1]
RP TISSUE=Bone marrow;
RC TISSUE=Bone marrow;
RX MEDLINE=96051398; PubMed=7584044;
RA Nomura N., Nagase T., Miyajima N., Sazuka T., Tanaka A., Sato S.,
RA Seki N., Kawabayashi Y., Ishikawa K.-I., Tabata S.;
RA "Prediction of the coding sequences of unidentified human genes. II.
RT The coding sequences of 40 new genes (KIAA0041-KIAA0080) deduced by

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RT analysis of cDNA clones from human cell line KG-1."
 RL DNA Res. 1:223-229(1994).
 RN [2]
 RP MEDLINE=98038990; PubMed=9373153;
 RX Thodpe K.L., Gorman P., Thomas C., Sheer D., Trowsdale J., Beck S.,
 RA "Chromosome 1 localization, gene structure and transcription pattern of
 RT the ORF gene, a homologue of the MHC-linked RING3 gene."
 RL Gene 200:177-183(1997).
 CC -1- SUBCELLULAR LOCATION: Nuclear (Potential).
 CC -1- TISSUE SPECIFICITY: Ubiquitous.
 CC -1- SIMILARITY: Contains 2 Bromodomains.
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 CC -----
 DR EMBL: D26362; BA05393.1; -
 DR EMBL: Z81330; CAB03630.1; -
 DR HSSP: Q92831; 1B91.
 DR Genew; HGNC:1104; BRD3.
 DR MIM: 601541; -
 DR GO: 0005634; C:nucleus; NAS.
 DR InterPro: IPR001487; Bromodomain.
 DR Pfam: PF00439; bromodomain; 2.
 DR PRINTS: PR00503; BROMODOMAIN.
 DR SMART; SM00297; BROMO; 2.
 DR PROSITE; PS00633; BROMODOMAIN 1; 2.
 DR PROSITE; PS50014; BROMODOMAIN 2; 2.
 DR Bromodomain; Repeat; Nuclear Protein.
 FT DOMAIN 56 115 BROMODOMAIN 1.
 FT DOMAIN 326 398 BROMODOMAIN 2.
 FT DOMAIN 487 555 LYS-RICH.
 FT DOMAIN 676 725 SER-RICH.
 FT CONFLICT 465 466 EL -> DV (IN REF. 2).
 FT SEQUENCE 726 AA; 79541 MW; 64F526FC3C1033AA CRC64;
 SQ
 Query Match 67.9%; Score 74; DB 1; Length 726;
 Best Local Similarity 70.0%; Pred. No. 0.42;
 Matches 14; Conservative 5; Mismatches 1; Indels 0; Gaps 0;
 Oy 2 KKKKKKKKKKKKKKKKKKK 21
 Db 489 KKKKKKKKKKKKKKKKKKK 508
 RESULT 11
 CBFS CANAL STANDARD; PRT; 479 AA.
 ID CBFS CANAL
 AC 043101;
 DT 15-JUL-1998 (Rel. 36; Created)
 DT 15-JUL-1998 (Rel. 36; Last sequence update)
 DT 28-FEB-2003 (Rel. 41; Last annotation update)
 DE Centromere/microtubule binding protein CBFS (Centromere-binding factor
 DE 5) (Nucleolar protein CBFS).
 GN CBFS.
 OS Candida albicans (Yeast).
 CC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 CC Saccharomycetales; mitosporic Saccharomycetales; Candida.
 CC NCB1_TaxID=5476;
 CC [1]
 CC SEQUENCE FROM N.A.
 RA Jiang W., Cliford J., Koltin Y.;
 RL Submitted (JAN1998) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: BINDS IN VITRO TO CENTROMERES AND MICROTUBULES. IT IS A
 CC CENTROMERE DNA-CBFS-BINDING FACTOR AND IS INVOLVED IN MITOTIC
 CC CHROMOSOME SEGREGATION. IT IS ESSENTIAL FOR CELL GROWTH. MAY BE IN
 CC SOME WAY ASSOCIATED WITH THE CBFS 110 KDA SUBUNIT (CBFS3A) (BY
 CC SIMILARITY).

CC -1- SUBCELLULAR LOCATION: Nuclear; nucleolar (by similarity).
 CC -1- SIMILARITY: BELONGS TO THE TRUB FAMILY OF PSEUDOURIDINE SYNTHASES.
 CC -1- SIMILARITY: Contains 1 PUA domain.
 CC -----
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 CC -----
 DR EMBL: U59149; AAB94297.1; -
 DR InterPro: IPR004802; Cof5.
 DR InterPro: IPR002478; PUA.
 DR InterPro: IPR002501; Trub_N.
 DR InterPro: IPR004521; Unchar_dom_2.
 DR Pfam; PF01472; PUA; 1.
 DR SMART; SM00359; PUA; 1.
 DR TIGRFAMs; TIGR00425; CBFS; 1.
 DR TIGRFAMs; TIGR00451; unchar_dom_2; 1.
 DR PROSITE; PS50890; PUA; 1.
 DR Microtubules; Centromere; Repeat; Nuclear protein; DNA-binding.
 KW Microtubules; Centromere; Repeat; Nuclear protein; DNA-binding.
 FT DOMAIN 267 342 PUA.
 FT SEQUENCE 479 AA; 54321 MW; 3BAF5104E12C9EB6 CRC64;
 SQ
 Query Match 66.1%; Score 72; DB 1; Length 479;
 Best Local Similarity 70.0%; Pred. No. 0.47;
 Matches 14; Conservative 4; Mismatches 2; Indels 0; Gaps 0;
 Oy 2 KKKKKKKKKKKKKKKKKKK 21
 Db 432 KKKKKKKKKKKKKKKKKKK 451
 RESULT 12
 GARP PLAFF STANDARD; PRT; 678 AA.
 ID GARP PLAFF
 AC P13816;
 DT 01-JAN-1990 (Rel. 13; Created)
 DT 01-JAN-1990 (Rel. 13; Last sequence update)
 DT 15-JUL-1999 (Rel. 38; Last annotation update)
 DE Glutamic acid-rich protein precursor.
 DE GARP.
 GN Plasmodium falciparum (Isolate FC27 / Papua New Guinea).
 CC Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.
 CC NCB1_TaxID=5837;
 CC [1]
 CC SEQUENCE FROM N.A.
 RX MEDLINE=89040048; PubMed=2903445;
 RA Trigila T., Stahl H.-D., Crewther P.E., Silva A., Anders R.F.,
 RA Kemp D.J.;
 RL "Structure of a Plasmodium falciparum gene that encodes a glutamic
 RL acid-rich protein (GARP)."
 RL Mol. Biochem. Parasitol. 31:199-202(1988).
 CC -----
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 CC -----
 DR EMBL: J03998; AAA29605.1; -
 DR PIR; A54514; A54514.
 KW Repeat; Malaria; Antigen; Signal.
 FT SIGNAL 1 25
 FT CHAIN 26 678 GLUTAMIC ACID-RICH PROTEIN.
 FT DOMAIN 120 164 15 X 3 AA TANDEN REPEATS OF K-K-X.
 FT DOMAIN 372 416 9 X APPROXIMATE TANDEN REPEATS.
 FT DOMAIN 417 441 5 X APPROXIMATE TANDEN REPEATS.


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FT DOMAIN 576 604 POLY-GLU.
FT DOMAIN 605 653 7 X APPROXIMATE TANDEM REPEATS.
FT DOMAIN 654 663 POLY-GLU.
SQ SEQUENCE 678 AA: 80551 MW: 2A8F856064962A9E CRC64;

Query Match 66.1%; Score 72; DB 1; Length 678;
Best Local Similarity 70.0%; Pred. No. 0.62;
Matches 14; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKK 21
DB 132 KKKKKKKKKKKKKKKKKKK 151

RESULT 13
CG79_HUMAN STANDARD; PRT; 351 AA.
AC Q9Y388;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DE 28-FEB-2003 (Rel. 41, Last annotation update)
DE Hypothetical protein CGI-79.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=20272150; PubMed=10810093;
RA Lai C.-H., Chou C.-Y., Ch'ang L.-Y., Liu C.-S., Lin W.-C.;
RT "Identification of novel human genes evolutionarily conserved in
RT Caenorhabditis elegans by comparative proteomics."
RL Genome Res. 10:703-713(2000).
CC -1- SIMILARITY: Contains 1 RNA recognition motif (RRM) domain.
CC -----
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CC -----
DR EMBL: AF151837; AAD34074.1; -.
DR HSSP: P26368; 2UZP.
DR InterPro: IPR00504; RNA_rec_mot.
DR Pfam: PF00076; rrm; 1.
DR SMART: SMO0360; RRM; 1.
DR PROSITE: PSS0102; RRM; 1.
DR PROSITE: PS00030; RRM_RNP_1; 1.
KM Hypothetical protein; RNA-binding.
FT DOMAIN 36 114 RNA-BINDING (RRM).
SQ SEQUENCE 351 AA: 39675 MW: 7B6E882D68192EBE CRC64;

Query Match 65.1%; Score 71; DB 1; Length 351;
Best Local Similarity 73.7%; Pred. No. 0.46;
Matches 14; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKK 20
DB 156 KKKKKKKKKKKKKKKKKKK 174

RESULT 14
CG1_HUMAN STANDARD; PRT; 686 AA.
AC P29973; Q16279; Q16485;
DT 01-APR-1993 (Rel. 25, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE cGMP-gated cation channel alpha 1 (CNG channel alpha 1) (CNG1)
DE (Cyclic nucleotide-gated cation channel alpha 1) (Cyclic nucleotide-gated
DE channel, photoreceptor) (Cyclic nucleotide-gated cation channel 1)

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DE (rod photoreceptor cGMP-gated channel alpha subunit).
GN CNG1 OR CNG1 OR CNGC.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Retina;
RX MEDLINE=92210603; PubMed=1372902;
RA Pittler S.J., Lee A.K., Altherr M.R., Howard T.A., Seldin M.F.,
RA Hurwitz R.L., Masmuch J.J., Baehr W.;
RT "Primary structure and chromosomal localization of human and mouse
RT rod photoreceptor cGMP-gated cation channel."
RL J. Biol. Chem. 267:6257-6262(1992).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Retina;
RX MEDLINE=92356211; PubMed=1379636;
RA Dhaliwal R.S., Macke J.P., Eddy R.L., Shows T.B., Reed R.R.,
RA Yau K.-W., Nathans J.;
RT "Human rod photoreceptor cGMP-gated channel: amino acid sequence,
RT gene structure, and functional expression."
RL J. Neurosci. 12:3248-3256(1992).
RN [3]
RP SEQUENCE OF 313-573 FROM N.A.
RX MEDLINE=95175019; PubMed=7532814;
RA Dietler W., Biel M., Flocke V., Hofmann F.;
RT "Expression of cyclic nucleotide-gated cation channels in non-sensory
RT tissues and cells."
RL Neuropharmacology 33:1275-1282(1994).
RN [4]
RP VARIANT ARR PHE-316, AND VARIANTS GLN-28 AND ASN-114.
RX MEDLINE=96036047; PubMed=7479749;
RA Dryja T.P., Finn J.T., Peng Y.-W., McGee T.L., Berson E.L., Yau K.-W.;
RT "Mutations in the gene encoding the alpha subunit of the rod
RT cGMP-gated channel in autosomal recessive retinitis pigmentosa."
RL Proc Natl. Acad. Sci. U.S.A. 92:10177-10181(1995).
CC -1- FUNCTION: VISUAL SIGNAL TRANSDUCTION IS MEDIATED BY A G-PROTEIN
CC COUPLED CASCADE USING GMP AS SECOND MESSENGER. THIS PROTEIN CAN
CC BE ACTIVATED BY CYCLIC GMP WHICH LEADS TO AN OPENING OF THE CATION
CC CHANNEL AND THEREBY CAUSING A DEPOLARIZATION OF ROD
CC PHOTORECEPTORS.
CC -1- SUBUNIT: HOMOTETRAMER OR HIGHER OLIGOMER. FORMS HETEROOLIGOMERIC
CC COMPLEX WITH CNG4.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein.
CC -1- TISSUE SPECIFICITY: ROD CELLS IN THE RETINA.
CC -1- DISEASE: Defects in CNG1 are a cause of autosomal recessive
CC retinitis pigmentosa (ARRP) [MIM:123825], a disease that leads to
CC degeneration of retinal photoreceptor cells.
CC -1- SIMILARITY: BELONGS TO THE CYCLIC NUCLEOTIDE-GATED CATION CHANNEL
CC FAMILY.
CC -1- DATABASE: NAME=Mutations of the CNG1 gene;
CC NOTE=Retina International's Scientific Newsletter;
CC WWW="http://www.retina-international.com/sci-news/cngalmut.htm".
CC -----
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CC -----
DR EMBL: M84741; AAA52010.1; ALT_INIT.
DR EMBL: S42457; AAB22778.1; -.
DR EMBL: S76062; AAD14206.1; -.
DR Genew: HGNC:2148; CNG1.
DR MIM: 123825; -.
DR GO: GO:0005887; C:integral to plasma membrane; TAS.
DR GO: GO:0006832; P:molecule transport; TAS.
DR GO: GO:0007601; P:vision; TAS.
DR InterPro: IPR000595; cGMP_binding.

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DR InterPro; IPR005821; Ion_trans.
DR InterPro; IPR001622; K+channel_pore.
DR Pfam; PF00027; CNMP_binding; 1.
DR Pfam; PF00520; IonTrans; 1.
DR SMART; SM00100; CNMP; 1.
DR PROSITE; PS00888; CNMP_BINDING_1; 1.
DR PROSITE; PS00889; CNMP_BINDING_2; 1.
DR PROSITE; PS00442; CNMP_BINDING_3; 1.
DR Ionic channel; Ion transport; CAMP-binding; Transmembrane;
KW Multigene family; Vision; Disease mutation; Polymorphism;
KW Retinitis pigmentosa.
FT DOMAIN 1 160
FT TRANSMEM 161 181
FT TRANSMEM 182 194
FT TRANSMEM 195 213
FT TRANSMEM 214 237
FT TRANSMEM 238 257
FT TRANSMEM 258 295
FT TRANSMEM 296 318
FT TRANSMEM 319 370
FT TRANSMEM 371 390
FT TRANSMEM 391 474
FT TRANSMEM 475 495
FT DOMAIN 496 686
FT NP_BIND 483 605
FT BINDING 542 542
FT BINDING 557 557
FT CARBOHYD 421 421
FT VARIANT 28 28
FT VARIANT 114 114
FT VARIANT 316 316
FT CONFLICT 46 46
FT CONFLICT 85 85
FT CONFLICT 146 147
FT CONFLICT 539 539
FT CONFLICT 677 678
SQ SEQUENCE 686 AA; 79126 MW; E5200D216FC97AF6 CRC64;

Query Match 65.1%; Score 71; DB 1; Length 686;
Best Local Similarity 70.0%; Pred. No. 0.78;
Matches 14; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKK 21
Db 124 KKKKKKKKKKKKKKKKKKK 143

RESULT 15
TCOF_HUMAN STANDARD; PRT; 1411 AA.
AC Q13428; Q99408; Q99860;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Treacle protein (Treacher Collins syndrome protein).
GN TCOF1.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP MEDLINE=96154183; PubMed=8563749;
RA Dixon J., Edwards S.J., Gladwin A.J., Dixon M.J., Loftus S.K.,
RA Bonner C.A., Koprivnikar K., Mammuth J.J.,
RT "Positional cloning of a gene involved in the pathogenesis of
RT Treacher Collins syndrome";
RL Nat. Genet. 12:130-136(1996).
RN [2]
RP SEQUENCE FROM N.A.

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RX MEDLINE=97228990; PubMed=9074926;
RA Dixon J., Edwards S.J., Anderson I., Brass A., Scambler P.J.,
RA Dixon M.J.;
RT "Identification of the complete coding sequence and genomic
RT organization of the Treacher Collins syndrome gene.";
RL Genome Res. 7:223-234(1997).
RN [3]
RP SEQUENCE FROM N.A.
RX MEDLINE=97250498; PubMed=9096354;
RA Wise C.A., Chiang L.C., Paznekas W.A., Sharma M., Musy M.M.,
RA Ashley J.A., Lovett M., Jabs E.W.;
RT "TCOF1 gene encodes a putative nuclear phosphoprotein that exhibits
RT mutations in Treacher Collins syndrome throughout its coding
RT region.";
RN [4]
RP VARIANTS LEU-439; VAL-810; VAL-1313 AND GLY-1355, AND VARIANT TCS
RP ARG-53.
RX MEDLINE=97195537; PubMed=9042910;
RA Edwards S.J., Gladwin A.J., Dixon M.J.;
RT "The mutational spectrum in Treacher Collins syndrome reveals a
RT predominance of mutations that create a premature-termination
RT codon.";
RL Am. J. Hum. Genet. 60:515-524(1997).
CC -1- DISEASE: Defects in TCOF1 are the cause of Treacher Collins
CC syndrome (TCS) [MIM:154500]. TCS is an autosomal dominant disorder
CC of craniofacial development that occurs with an incidence of
CC 1/50,000 live births. The clinical features of TCS are bilaterally
CC symmetrical and include: (1) abnormalities of the external ears,
CC atresia of the external ear canals, and malformation of the middle
CC ear ossicles, which may result in conductive hearing loss; (2)
CC lateral downward sloping of palpebral fissures, frequently with
CC colobomas of the lower eyelids; (3) hypoplasia of the mandible and
CC zygomatic complex; (4) cleft palate.
CC -1- SIMILARITY: Contains 1 Lish domain.
CC -----
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DR EMBL; U40847; AAC50903.1; -
DR EMBL; U76366; AAC51181.1; -
DR EMBL; U84664; AAC51185.1; JOINED.
DR EMBL; U84640; AAC51185.1; JOINED.
DR EMBL; U84641; AAC51185.1; JOINED.
DR EMBL; U84642; AAC51185.1; JOINED.
DR EMBL; U84643; AAC51185.1; JOINED.
DR EMBL; U84644; AAC51185.1; JOINED.
DR EMBL; U84645; AAC51185.1; JOINED.
DR EMBL; U84646; AAC51185.1; JOINED.
DR EMBL; U84647; AAC51185.1; JOINED.
DR EMBL; U84648; AAC51185.1; JOINED.
DR EMBL; U84649; AAC51185.1; JOINED.
DR EMBL; U84650; AAC51185.1; JOINED.
DR EMBL; U84651; AAC51185.1; JOINED.
DR EMBL; U84652; AAC51185.1; JOINED.
DR EMBL; U84653; AAC51185.1; JOINED.
DR EMBL; U84654; AAC51185.1; JOINED.
DR EMBL; U84655; AAC51185.1; JOINED.
DR EMBL; U84656; AAC51185.1; JOINED.
DR EMBL; U84657; AAC51185.1; JOINED.
DR EMBL; U84658; AAC51185.1; JOINED.
DR EMBL; U84659; AAC51185.1; JOINED.
DR EMBL; U84660; AAC51185.1; JOINED.
DR EMBL; U84661; AAC51185.1; JOINED.
DR EMBL; U84662; AAC51185.1; JOINED.
DR EMBL; U84663; AAC51185.1; JOINED.
DR EMBL; U79659; AAB40722.1; -
DR EMBL; U79645; AAB40722.1; JOINED.

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DR EMBL; U79646; AAB40722.1; JOINED.
DR EMBL; U79647; AAB40722.1; JOINED.
DR EMBL; U79648; AAB40722.1; JOINED.
DR EMBL; U79649; AAB40722.1; JOINED.
DR EMBL; U79650; AAB40722.1; JOINED.
DR EMBL; U79651; AAB40722.1; JOINED.
DR EMBL; U79652; AAB40722.1; JOINED.
DR EMBL; U79653; AAB40722.1; JOINED.
DR EMBL; U79654; AAB40722.1; JOINED.
DR EMBL; U79655; AAB40722.1; JOINED.
DR EMBL; U79656; AAB40722.1; JOINED.
DR EMBL; U79657; AAB40722.1; JOINED.
DR EMBL; U79658; AAB40722.1; JOINED.
DR Genew; HGNC:11654; TCOF1.
DR MIM; 606847; -.
DR MIM; 154500; -.
DR GO; GO:0005730; C:nucleolus; TAS.
DR GO; GO:0005215; F:transporter activity; TAS.
DR GO; GO:0001501; P:skeletal development; TAS.
DR InterPro; IPR006594; LISH.
DR InterPro; IPR003993; treacle.
DR Pfam; PF03546; treacle; 3.
DR PRINTS; PR01503; TREACLE.
DR SMART; SM00667; LISH; 1.
DR PROSITE; PS00896; LISH; 1.
KW Disease mutation; Polymorphism.
FT DOMAIN 6 38 LISH.
FT DOMAIN 89 97 POLY-GLU.
FT DOMAIN 204 207 POLY-SER.
FT DOMAIN 616 619 POLY-SER.
FT DOMAIN 919 924 POLY-SER.
FT DOMAIN 1285 1289 POLY-LYS.
FT DOMAIN 1375 1386 POLY-LYS.
FT DOMAIN 1398 1405 POLY-LYS.
FT VARIANT 53 53 W -> R (in TCS).
FT VARIANT 439 439 /FTid=VAR_005630.
FT VARIANT 810 810 P -> L.
FT VARIANT 810 810 /FTid=VAR_005631.
FT VARIANT 1313 1313 A -> V.
FT VARIANT 1355 1355 /FTid=VAR_005632.
FT VARIANT 1355 1355 A -> V (in dbSNP:15251).
FT CONFLICT 1312 1312 D -> G.
FT CONFLICT 1411 AA; 144312 MM; 3880203D985C2699 CRC64;
SQ SEQUENCE 1411 AA; 144312 MM; 3880203D985C2699 CRC64;

Query Match 64.7%; Score 70.5; DB 1; Length 1411;
Best Local Similarity 54.8%; Pred. No. 1.5;
Matches 17; Conservative 2; Mismatches 1; Indels 11; Gaps 1;

Qy 2 KKKKKKKKKKK-----KKKKKKK 21
Db 1375 KEKKEKKKKAKKASTKDSKSPQKKKKKKK 1405
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Search completed: January 30, 2004, 00:20:44
Job time : 5.73239 secs

OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 OC Eriocaridaceae; Oryzae; Oryza.
 OK NCBI_TaxID=4530;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Nipponbare;
 RA Buell C.R., Yuan Q., Ouyang S., Liu J., Moffat K.S., Hill J.N.,
 RA Ganabinger K., Brenner M., Burgess S., Hance M., Shvartsbeyn M.,
 RA Tselirin T., Riggs F., Hsiao J., Zismann V., Blunt S., Pal G.,
 RA Vanhaken S.E., Uteback T.R., Feldblyum T.V., Kalb E., Quackenbush J.,
 RA Salzberg S.L., White O., Fraser C.M.;
 RT "Oryza sativa chromosome 10 BAC OSJNBa0057L21 genomic sequence.";
 RL Submitted (FEB-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AC087599; AAL79706.1; -.
 DR Gramene; Q857D3; -.
 KW Hypothetical protein.
 SQ SEQUENCE 80 AA; 9362 MW; 017C86313B21D8 CRC64;

Query Match 91.7%; Score 100; DB 10; Length 80;
 Best Local Similarity 100.0%; Pred. No. 0.0001;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKK 21
 DB 48 KKKKKKKKKKKKKKKKKKK 67

RESULT 3
 Q8LOP6 PRELIMINARY; PRT; 113 AA.

DT 01-OCT-2002 (TREMBlrel. 22, Created)
 DT 01-OCT-2002 (TREMBlrel. 22, Last sequence update)
 DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
 DE OJ117_G01.13 protein.
 GN OJ117_G01.13.
 OS Oryza sativa (japonica cultivar-group).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 OC Eriocaridaceae; Oryzae; Oryza.
 OK NCBI_TaxID=39947;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=cv. Nipponbare;
 RA Sasaki T., Matsunoto T., Yamamoto K.;
 RT "Oryza sativa nipponbare (GA3) genomic DNA, chromosome 1, BAC
 RT clone:OJ117_G01.";
 RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AP003374; BAB93330.1; -.
 DR Gramene; Q8LOP6; -.
 SQ SEQUENCE 113 AA; 13660 MW; 597DB0E8E2MA33EF CRC64;

Query Match 91.7%; Score 100; DB 10; Length 113;
 Best Local Similarity 100.0%; Pred. No. 0.00013;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKK 21
 DB 10 KKKKKKKKKKKKKKKKKKK 29

RESULT 4
 Q9P529 PRELIMINARY; PRT; 128 AA.

DT 01-OCT-2000 (TREMBlrel. 15, Created)
 DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
 DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
 DE Hypothetical 15.2 kDa protein.
 GN B24H17.160.
 OS Neurospora crassa.
 OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;

OC Sordariales; Sordariaceae; Neurospora.
 OK NCBI_TaxID=5141;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Schulte U., Aign V., Heiseisel J., Brandt P., Fartmann B., Holland R.,
 RA Nyakatura G., Mewes H.W., Mannhaupt G.;
 RL Submitted (MAY-2000) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA German Neurospora genome project;
 RL Submitted (OCT-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AL356815; CAB92638.2; -.
 KW Hypothetical protein.
 SQ SEQUENCE 128 AA; 15157 MW; 8C7C65C3DFB70765 CRC64;

Query Match 91.7%; Score 100; DB 3; Length 128;
 Best Local Similarity 100.0%; Pred. No. 0.00014;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKK 21
 DB 71 KKKKKKKKKKKKKKKKKKK 90

RESULT 5
 O35807 PRELIMINARY; PRT; 129 AA.

DT 01-JAN-1998 (TREMBlrel. 05, Created)
 DT 01-JAN-1998 (TREMBlrel. 05, Last sequence update)
 DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
 DE MICROVASCULAR endothelial differentiation protein 2.
 GN MDG2.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OK NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISUB=Epilidymis;
 RX MEDLINE=98172708; PubMed=9511718;
 RA Proels F., Loser B., Marx M.;
 RT "Differential expression of osteopontin, PC4, and CEC5, a novel mRNA
 RT species, during in vitro angiogenesis.";
 RL Exp. Cell Res. 239:1-10(1998).
 DR EMBL; Y08769; CAA70022.1; -.
 DR InterPro; IPR000719; Prot_Kinase.
 DR Pfam; PF00069; Kinase; 1.
 DR ProDom; PD000001; Prot_kinase; 1.
 DR PROSITE; PS50011; PROTEIN_KINASE_DOM; 1.
 KW ATP-binding; Transferase.
 SQ SEQUENCE 129 AA; 15080 MW; 38102272B8E2EDB4 CRC64;

Query Match 91.7%; Score 100; DB 11; Length 129;
 Best Local Similarity 100.0%; Pred. No. 0.00014;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKK 21
 DB 85 KKKKKKKKKKKKKKKKKKK 104

RESULT 6
 Q9H5V6 PRELIMINARY; PRT; 168 AA.

DT 01-MAR-2001 (TREMBlrel. 16, Created)
 DT 01-MAR-2001 (TREMBlrel. 16, Last sequence update)
 DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
 DE Hypothetical protein FLJ22976 (Fragment).
 GN Homo sapiens (Human).
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

```

OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Watanabe K., Kunagai A., Itakura S., Yamazaki M., Tashiro H., Ota T.,
RA Suzuki Y., Ogasashi M., Nishi T., Shibahara T., Tanaka T.,
RA Nakamura Y., Isogai T., Sugano S.;
RT "NEDO human cDNA sequencing project.";
RL Submitted (Aug-2000) to the EMBL/Genbank/DBJ databases.
DR EMBL; AK026629; BAB15513.1; -
KW Hypothetical protein.
FT NON TER 168
SQ SEQUENCE 168 AA; 19549 MW; A19DBD195FB8A1A90 CRC64;

Query Match
Best Local Similarity 91.7%; Score 100; DB 4; Length 168;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 KKKKKKKKKKKKKKKKKKKKK 21
Db 140 KKKKKKKKKKKKKKKKKKKKK 159

RESULT 7
O81247 PRELIMINARY; PRT; 206 AA.
AC O81247;
DT 01-MAR-2003 (TReMBLrel. 23, Created)
DT 01-MAR-2003 (TReMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TReMBLrel. 23, Last annotation update)
DE Hypothetical protein.
PF0475C.
OS Plasmodium falciparum (isolate 3D7).
OC Eukaryota; Alveolata; Apicomplexa; Hemosporidia; Plasmodium.
OX NCBI_TaxID=36329;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22255708; PubMed=12368867;
RA Hall N., Pain A., Berriman M., Churcher C., Harris B., Harris D.,
RA Mungall K., Bowman S., Atkin R., Baker S., Barron A., Brooks K.,
RA Buckee C.O., Burrows C., Cherevach I., Chillingworth C.,
RA Chillingworth T., Christodoulou Z., Clark L., Clark R., Corton C.,
RA Cronin A., Davies R., Davis P., Dear P., Dearden F., Doggett J.,
RA Feltwell T., Goble A., Goodhead I., Gwilliam R., Hamlin N., Hance Z.,
RA Harper D., Hauser H., Hornsby T., Holtroyd S., Horrocks P.,
RA Humphrey S., Jagsels K., James K.D., Johnson D., Kerhornou A.,
RA Knighes A., Konfortov B., Kyes S., Larke N., Lawson D., Lemard N.,
RA Line A., Maddison M., McLean J., Mooney P., Moule S., Murphy L.,
RA Oliver K., Ormond D., Price C., Quail M.A., Rabinowitz E.,
RA Rajandream M.A., Rutter S., Rutherford K.M., Sanders M., Simmonds M.,
RA Seeger K., Sharp S., Smith R., Squares R., Stevens K.,
RA Taylor K., Tivey A., Unwin L., Whitehead S., Woodward J.,
RA Sulston J.E., Craig A., Newbold C., Barrell B.G.;
RT "Sequence of Plasmodium falciparum chromosomes 1, 3-9 and 13.";
RL Nature 419:527-531(2002).
DR EMBL; AL031745; CAD49055.1; -
KW Hypothetical protein.
SQ SEQUENCE 206 AA; 25047 MW; 1192E49A3DC4523F CRC64;

Query Match
Best Local Similarity 91.7%; Score 100; DB 5; Length 206;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 KKKKKKKKKKKKKKKKKKKKK 21
Db 185 KKKKKKKKKKKKKKKKKKKKK 204

RESULT 8
O64075 PRELIMINARY; PRT; 215 AA.
AC O64075;
DT 01-NOV-1996 (TReMBLrel. 01, Created)
DT 01-NOV-1996 (TReMBLrel. 01, Last sequence update)

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DT 01-OCT-2002 (TReMBLrel. 22, Last annotation update)
DE Nucleoporin p62 homolog protein (Fragment).
OS Rattus sp.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10118;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=95151924; PubMed=7849178;
RA Wang Z.Q., Akmal K.M., Kim K.H.;
RT "An unusual nucleoporin-related messenger ribonucleic acid is present
RT in the germ cells of rat testis.";
RL Biol. Reprod. 51:1022-1030(1994).
DR EMBL; S75997; AAB33384.1; -
KW Porin.
FT NON TER 1
SQ SEQUENCE 215 AA; 24593 MW; 098251C97A8FB88 CRC64;

Query Match
Best Local Similarity 91.7%; Score 100; DB 11; Length 215;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 KKKKKKKKKKKKKKKKKKKKK 21
Db 35 KKKKKKKKKKKKKKKKKKKKK 54

RESULT 9
O9LG29 PRELIMINARY; PRT; 260 AA.
AC O9LG29;
DT 01-OCT-2000 (TReMBLrel. 15, Created)
DT 01-OCT-2000 (TReMBLrel. 15, Last sequence update)
DT 01-OCT-2002 (TReMBLrel. 22, Last annotation update)
DE Genomic DNA, chromosome 3, BAC clone:FLD9.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC Eustoids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Colombia;
RA Nakamura Y.;
RT "Structural Analysis of Arabidopsis thaliana Chromosome 3. III.";
RL Submitted (JUN-2000) to the EMBL/Genbank/DBJ databases.
DR EMBL; AP002460; BAA97098.1; -
DR InterPro; IPR005819; Histone_H5.
DR PRINTS; PR00624; HISTONEH5.
SQ SEQUENCE 260 AA; 33307 MW; 43E2394CB8131143 CRC64;

Query Match
Best Local Similarity 91.7%; Score 100; DB 10; Length 260;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 KKKKKKKKKKKKKKKKKKKKK 21
Db 7 KKKKKKKKKKKKKKKKKKKKK 26

RESULT 10
O9NT34 PRELIMINARY; PRT; 380 AA.
AC O9NT34;
DT 01-OCT-2000 (TReMBLrel. 15, Created)
DT 01-OCT-2000 (TReMBLrel. 15, Last sequence update)
DT 01-OCT-2002 (TReMBLrel. 22, Last annotation update)
DE Hypothetical protein (Fragment).
GN DKFZP434I1120.
OS Homo sapiens (human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;

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RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Testis;
RA Otterweiler B., Obermaier B., Mewes H.W., Gassenhuber J., Wiemann S.;
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AL1137556; CAB70810.1; -
DR Genew; HGNC:15736; C17orf28.
KW Hypothetical protein.
FT NON TER 380
SQ SEQUENCE 380 AA; 42689 MW; 67F50DD101346AFB CRC64;

Query Match 91.7%; Score 100; DB 4; Length 380;
Best Local Similarity 100.0%; Pred. No. 0.00028;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKKKK 21
DB 355 KKKKKKKKKKKKKKKKKKK 374

RESULT 11
ID Q8SWR7 PRELIMINARY; PRT; 515 AA.
AC Q8SWR7;
DT 01-JUN-2002 (TREMBlrel. 21, Created)
DT 01-JUN-2002 (TREMBlrel. 21, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE GH22607p (Fragment).
GN CG7180.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neuroptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
CX NCBI_Taxid=7227;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Berkley;
RA Stapleton M., Brokstein P., Hong L., Aspayani A., Carlson J.,
RA Champagne M., Chavez C., Dorsett V., Drensek D., Farfan D., Frise E.,
RA George R., Gonzalez M., Guarin H., Krommiller B., Li P., Liao G.,
RA Miranda A., Mungall C.J., Nuno J., Pacleb J., Paragas V., Park S.,
RA Patel S., Phouanavong S., Wan K., Yu C., Lewis S.E., Rubin G.M.,
RA Ceiniker S.;
RL Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY055518; AAM12251.1; -
DR FlyBase; FBgn0032673; CG7180.
DR InterPro; IPR000387; TYR_phosphatase.
DR Pfam; PF00102; Y_phosphatase; 1.
DR SMART; SM00194; PTPc; 1.
DR PROSITE; PS00383; TYR_PHOSPHATASE_1; 1.
DR PROSITE; PS50056; TYR_PHOSPHATASE_2; 1.
DR PROSITE; PS50055; TYR_PHOSPHATASE_PTP; 1.
KW Hydroxylase.
FT NON TER 515
SQ SEQUENCE 515 AA; 59080 MW; B2825B7BEA96195E CRC64;

Query Match 91.7%; Score 100; DB 5; Length 515;
Best Local Similarity 100.0%; Pred. No. 0.00034;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKKKK 21
DB 493 KKKKKKKKKKKKKKKKKKK 512

RESULT 12
ID Q9LXR2 PRELIMINARY; PRT; 517 AA.
AC Q9LXR2;
DT 01-OCT-2000 (TREMBlrel. 15, Created)
DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)

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DE Hypothetical 59.7 kDa protein.
GN T20N10.250.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
CX NCBI_Taxid=3702;
RN [1]
RP SEQUENCE FROM N.A.
RA D'Angelo M., Verzi A., Modesto D., Pigazzi M., Valle G., Mewes H.W.,
RA Rudd S., Lemcke K., Mayer K.F.X., Queller F., Salanoubat M.;
RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
RP EU Arabidopsis sequencing project;
RA Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
RL EMBL; ALJ53032; CAB88307.1; -
DR InterPro; IPR001810; F-box.
DR Pfam; PF00646; F-box; 1.
DR SMART; SM00579; FBD; 1.
DR PROSITE; PS50181; FBOX; 1.
KW Hypothetical protein.
SQ SEQUENCE 517 AA; 59689 MW; EC6D957D01F86E70 CRC64;

Query Match 91.7%; Score 100; DB 10; Length 517;
Best Local Similarity 100.0%; Pred. No. 0.00034;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKKKK 21
DB 444 KKKKKKKKKKKKKKKKKKK 463

RESULT 13
ID Q95LV6 PRELIMINARY; PRT; 531 AA.
AC Q95LV6;
DT 01-DEC-2001 (TREMBlrel. 19, Created)
DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
DE Hypothetical 61.4 kDa protein (Fragment).
OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
OC Cercopithecinae; Macaca.
CX NCBI_Taxid=9541;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Testis;
RA Hashimoto K., Oseada N., Hida M., Kueuda J., Tanuma R., Hirai M.,
RA Terao K., Sugano S.;
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB071085; BAB64479.1; -
KW Hypothetical protein.
FT NON TER 531
SQ SEQUENCE 531 AA; 61389 MW; B55996B4F5CD60C CRC64;

Query Match 91.7%; Score 100; DB 6; Length 531;
Best Local Similarity 100.0%; Pred. No. 0.00035;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 KKKKKKKKKKKKKKKKKKKKK 21
DB 502 KKKKKKKKKKKKKKKKKKK 521

RESULT 14
ID Q9H6Q7 PRELIMINARY; PRT; 720 AA.
AC Q9H6Q7;
DT 01-OCT-2000 (TREMBlrel. 15, Created)
DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)

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AC Q9H6Q7;
 DT 01-MAR-2001 (TREMBLrel. 16, Created)
 DT 01-MAR-2001 (TREMBLrel. 16, last sequence update)
 DT 01-OCT-2002 (TREMBLrel. 22, last annotation update)
 DE Hypothetical protein FLJ21979 (Fragment).
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Kawabata A., Hiki J. T., Kobatake N., Inagaki H., Ikema Y., Okamoto S.,
 RA Okitani R., Ota T., Suzuki Y., Obayashi M., Nishi T., Shibahara T.,
 RA Tanaka T., Nakamura Y., Isogai T., Sugano S.;
 RT "NEDO human cDNA sequencing project."
 RL Submitted (Aug-2000) to the EMBL/Genbank/DBJ databases.
 DR EMBL; AK025632; BAB15196.1; -.
 KW Hypothetical protein.
 FT NON TER 720
 SQ SEQUENCE 720 AA; 84029 MW; A86586FEA953D0B CRC64;

Query Match 91.7%; Score 100; DB 4; Length 720;
 Best Local Similarity 100.0%; Pred. No. 0.00043;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 KKKKKKKKKKKKKKKKKKKKK 21
 DB 692 KKKKKKKKKKKKKKKKKKKKK 711

RESULT 15

ID O8T2U7 PRELIMINARY; PRT; 791 AA.
 AC O8T2U7;
 DT 01-JUN-2002 (TREMBLrel. 21, Created)
 DT 01-JUN-2002 (TREMBLrel. 21, last sequence update)
 DT 01-MAR-2003 (TREMBLrel. 23, last annotation update)
 DE Hypothetical 92.4 kDa protein.
 OS Dictyostelium discoideum (slime mold).
 OC Eukaryota; Mycetozoa; Dictyostelida; Dictyostelium.
 OX NCBI_TaxID=44689;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=AX4;
 RA Gloeckner G., Eichinger L., Szafranski K., Pachebat J., Dear P.,
 RA Lehmann R., Baumgart C., Parra G., April J.F., Guigo R., Kumpf K.,
 RA Tunggal B., Cox B., Quail M.A., Platzer M., Rosenthal A., Noegel A.A.;
 RT "Sequence and Analysis of Chromosome 2 of Dictyostelium."
 RL Submitted (MAR-2002) to the EMBL/Genbank/DBJ databases.
 DR EMBL; AC115574; AAL92183.1; -.
 DR InterPro; IPR005033; YEATS.
 DR InterPro; IPR007087; ZnF_C2H2.
 DR Pfam; PF03366; YEATS; 1.
 DR SMART; SM00355; ZnF_C2H2; 1.
 DR SMART; SM00355; ZnF_C2H2; 1.
 KW Hypothetical protein.
 SQ SEQUENCE 791 AA; 92375 MW; D66C6BDEC92352C CRC64;

Query Match 91.7%; Score 100; DB 5; Length 791;
 Best Local Similarity 100.0%; Pred. No. 0.00045;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 KKKKKKKKKKKKKKKKKKKKK 21
 DB 769 KKKKKKKKKKKKKKKKKKKKK 788

Search completed: January 30, 2004, 00:24:37
 Job time : 22.6901 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 30, 2004, 00:13:12 ; Search time 10.6808 Seconds
(without alignments)
225.098 Million cell updates/sec

Title: US-09-461-684C-2
Perfect score: 109
Sequence: 1 CEAANAAEAANAAEAANAAEAANAA 25

Scoring table:
BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283308 seqs, 9616862 residues
Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	77	70.6	265	2	S19113
2	72	66.1	165	2	B87702
3	71	65.1	205	2	S19114
4	70	64.2	183	2	S24960
5	70	64.2	401	2	A48423
6	69	63.3	97	2	S02376
7	69	63.3	1028	2	A56038
8	69	63.3	1213	2	S16356
9	68	63.3	1668	2	T11748
10	68	62.4	403	2	A81882
11	68	62.4	655	2	A29945
12	68	62.4	873	2	B53225
13	68	62.4	2023	2	T13154
14	67	61.5	314	2	UC5273
15	67	61.5	460	2	TJ3110
16	67	61.5	494	2	A42170
17	67	61.5	497	2	JC5076
18	67	61.5	604	2	A33369
19	67	61.5	606	2	S13367
20	67	61.5	627	2	T02610
21	67	61.5	671	2	C96534
22	67	61.5	1226	2	T24045
23	67	61.5	2715	2	T13049
24	66.5	60.1	543	2	B33369
25	65.5	60.1	273	2	T51010
26	65	59.6	85	1	A22592
27	65	59.6	333	2	A39065
28	65	59.6	475	2	A43915
29	65	59.6	475	2	A43915

30	65	59.6	644	2	S39356	transcription fact
31	65	59.6	703	2	T48600	kinase-like protei
32	64	58.7	109	1	R48P1	acidic ribosomal p
33	64	58.7	392	2	B48423	homeotic protein e
34	64	58.7	1065	2	T13230	dachshund isoform
35	64	58.7	1072	2	T13232	dachshund protein
36	64	58.7	1074	2	T13229	dachshund protein
37	64	58.7	1081	2	T13231	abdominal segment
38	64	58.7	1533	2	A46221	female sterile hom
39	64	58.7	2038	2	A43742	50S ribosomal prot
40	63	57.8	179	2	AF2908	polyomavirus enhan
41	63	57.8	179	2	F97683	asparaginyl-tna s
42	63	57.8	513	2	A48233	conserved hypocher
43	63	57.8	568	2	T39675	GTP-binding regula
44	63	57.8	581	2	E75383	
45	63	57.8	846	2	S52418	

ALIGNMENTS

```

RESULT 1
S19113
cgcr-4 protein - Chlamydomonas reinhardtii (fragment)
C:Species: Chlamydomonas reinhardtii
C>Date: 13-Jan-1995 #sequence_revision 13-Jan-1995 #text_change 21-Jul-2000
C:Accession: S19113; S14466
R:Wakarchuk, W.W.; Mueller, F.W.; Beck, C.F.
Plant Mol. Biol. 18, 143-146, 1992
A:Title: Two GC-rich DNA elements of Chlamydomonas reinhardtii with complex arrangements
A:Reference number: S19113; MUID:92119224; PMID:1731966
A:Accession: S19113
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-265 <MAX>
A:Cross-references: EMBL:X17208; NID:g18136; PIDN:CAA35080.1; PID:g18137
C:Genetics:
A:Gene: cgcr-4

Query Match      70.6%  Score 77;  DB 2;  Length 265;
Best Local Similarity 82.6%  Pred. No. 0.25;
Matches 19;  Conservative 1;  Mismatches 3;  Indels 0;  Gaps 0;

Cy      3 AAAAAAAAAAAAAAAAAAAAAA 25
Db      154 AAAAAAAAAAAKARVAAEARRAA 176

RESULT 2
B87702
ribosomal protein S16 [imported] - Caulobacter crescentus
C:Species: Caulobacter crescentus
C>Date: 20-Apr-2001 #sequence_revision 20-Apr-2001 #text_change 20-Apr-2001
C:Accession: B87702
R:Nierman, W.C.; Feldblum, T.V.; Paulsen, I.T.; Nelson, K.E.; Bisen, J.; Heidelberg, J.;
B.; Laub, M.T.; DeBoy, R.T.; Dodson, R.J.; Durkin, A.S.; Gwinn, M.L.; Haft, D.H.; Kolon
n, J.; Smolova, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Venter, J.C.; Fraser, C.M.
Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001
A:Title: Complete Genome Sequence of Caulobacter crescentus.
A:Reference number: A87249; MUID:21173698; PMID:11259647
A:Accession: B87702
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-165 <STO>
A:Cross-references: GB:AE005673; NID:g13425408; PIDN:AAK25614.1; GSPDB:GN00148
C:Genetics:
A:Gene: CC3652

Query Match      66.1%  Score 72;  DB 2;  Length 165;
Best Local Similarity 75.0%  Pred. No. 0.56;
Matches 18;  Conservative 2;  Mismatches 4;  Indels 0;  Gaps 0;

Cy      2 EAAAAAAAAAAAAAAAAAAAAA 25

```

DB 115 QAEADAKAAAEAKAAAEAAAAA 138

RESULT 3

S19114
C:Spec: Chlamydomonas reinhardtii (fragment)
C>Date: 13-Jan-1995 #sequence_revision 13-Jan-1995 #text_change 03-Nov-2000
C:Accession: S19114
R:Wakarchuk, W.W.; Mueller, F.W.; Beck, C.F.
Plant Mol. Biol. 18, 143-146, 1992
A>Title: Two GC-rich DNA elements of Chlamydomonas reinhardtii with complex arrangements
A:Reference number: S19113; MUID:92119224; PMID:1731966
A:Accession: S19114
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-205 <WAK>
A:Cross-references: EMBL:X17207
C:Superfamily: phage lambda hypothetical protein 401

Query Match 65.1%; Score 71; DB 2; Length 205;
Best Local Similarity 75.0%; Pred. No. 0.82;
Matches 18; Conservative 1; Mismatches 5; Indels 0; Gaps 0;

QY 2 EAAAAEAAAAEAAAAEAAAAA 25
DB 49 EAAAAEAAAAEAAAAEAAAAA 72

RESULT 4

S24960
gene C98 protein - rape
C:Species: Brassica napus (rape)
C>Date: 20-Feb-1995 #sequence_revision 20-Feb-1995 #text_change 09-Sep-1997
C:Accession: S24960
R:Robert, M.R.; Hodges, R.; Sorensen, A.; Ross, J.; Murphy, D.J.; Draper, R.; Scott, R.
submitted to the EMBL Data Library, July 1992
A:Description: A new class of Brassica napus oleosin genes specific to the male gametoph
A:Reference number: S24960
A:Accession: S24960
A>Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-183 <ROB>
A:Cross-references: EMBL:X67142; NID:g17792; PID:g17793

Query Match 64.2%; Score 70; DB 2; Length 183;
Best Local Similarity 78.3%; Pred. No. 0.95;
Matches 18; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 3 AAAAAEAAAAEAAAAEAAAAA 25
DB 151 AAPAAEPAAEAPAAEAPAA 173

RESULT 5

A48423
engrailed homeodomain-containing protein En-1 - mouse
N:Alternate names: homeotic protein En-1
C:Species: Mus musculus (house mouse)
C>Date: 01-Dec-1993 #sequence_revision 18-Nov-1994 #text_change 24-Sep-1999
C:Accession: A48423; S13009; A26629; A24778
R:Logan, C.; Hanks, M.C.; Noble-Topham, S.; Nallainathan, D.; Provar, N.J.; Joyner, A.L.
Dev. Genet. 13, 345-358, 1992
A>Title: Cloning and sequence comparison of the mouse, human, and chicken engrailed gene
A:Reference number: A48423; MUID:93185339; PMID:1363401
A:Accession: A48423
A>Status: preliminary; not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-401 <LOG>
A:Experimental source: CD-1, embryo
A>Note: sequence extracted from NCBI backbone (NCBIP:126620)
R:Holland, P.W.H.; Williams, N.A.

FEBS Lett. 277, 250-252, 1990
A>Title: Conservation of engrailed-like homeobox sequences during vertebrate evolution.
A:Reference number: S13009; MUID:91099509; PMID:1980115
A:Accession: S13009
A>Status: preliminary
A:Molecule type: nucleic acid
A:Residues: 321-380 <HOL>
R:Joyner, A.L.; Martin, G.R.
Genes Dev. 1, 29-38, 1987

A>Title: En-1 and En-2, two mouse genes with sequence homology to the Drosophila engrail
A:Reference number: A91620; MUID:88112776; PMID:2892757
A:Accession: A26629
A:Molecule type: DNA; mRNA
A:Residues: 278-401 <JOY>
A:Cross-references: GB:Y00201; GB:M11987; NID:g49587; PID:CA68361.1; PID:g669105
R:Joyner, A.L.; Kornberg, T.; Coleman, K.G.; Cox, D.R.; Martin, G.R.
Cell 43, 29-37, 1985

A>Title: Expression during embryogenesis of a mouse gene with sequence homology to the D
A:Reference number: A24778; MUID:86079501; PMID:2416459
A:Accession: A24778
A:Molecule type: DNA
A:Residues: 311-401 <JO2>
C:Genetics:
A:Gene: en.1
A:Map position: 1

C:Superfamily: unassigned homeobox proteins; homeobox homology
C:Keywords: DNA binding; homeobox; nucleus; transcription regulation
F:313-369/Domain: homeobox homology <HOX>

Query Match 64.2%; Score 70; DB 2; Length 401;
Best Local Similarity 78.3%; Pred. No. 1.7;
Matches 18; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 3 AAAAAEAAAAEAAAAEAAAAA 25
DB 207 AAAAAEAAAAEAAAAEAAAAA 229

RESULT 6

S02376
antifreeze protein precursor - yellowtail flounder
C:Species: Limanda ferruginea (yellowtail flounder)
C>Date: 01-Dec-1989 #sequence_revision 01-Dec-1989 #text_change 24-Oct-2000
C:Accession: S02376
R:Scott, G.K.; Davies, P.L.; Shears, M.A.; Fletcher, G.L.
Eur. J. Biochem. 168, 629-633, 1987

A>Title: Structural variations in the alanine-rich antifreeze proteins of the Pleuronecti
A:Reference number: S02376; MUID:88029483; PMID:3665937
A:Accession: S02376
A:Molecule type: mRNA
A:Residues: 1-97 <SCO>

A:Cross-references: EMBL:X06356; NID:g64041; PID:CAA29655.1; PID:g64042
A>Note: part of this sequence, including the amino end of the mature protein, was confirm
C:Superfamily: antifreeze protein
C:Keywords: antifreeze

F:1-23/Domain: signal sequence #status predicted <SIG>
F:24-48/Domain: propeptide #status predicted <PRO>
F:49-96/Product: antifreeze protein #status predicted <MAT>

Query Match 63.3%; Score 69; DB 2; Length 97;
Best Local Similarity 70.8%; Pred. No. 0.75;
Matches 17; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

QY 2 EAAAAEAAAAEAAAAEAAAAA 25
DB 53 DAAAAAATTAATAAAKAAADTAA 76

RESULT 7

A56038
DNA-binding protein ovo - fruit fly (Drosophila melanogaster)
C:Species: Drosophila melanogaster
C>Date: 01-Dec-1995 #sequence_revision 01-Dec-1995 #text_change 21-Jul-2000

C/Accession: A56038
R/Gartinkel, M.D.; Wang, J.; Mahowald, A.P.
Mol. Cell. Biol. 14:6809-6818, 1994
A/Title: Multiple products from the shavenbaby-ovo gene region of Drosophila melanogaster
A/Reference number: A56038; MUID:95021209; PMID:7935398
A/Accession: A56038
A/Status: preliminary
A/Molecule type: mRNA
A/Residues: 1-1028 <GAR>
A/Cross-references: GB:U1183; NID:9520526; PIDN:AA60216.1; PID:9520527
C/Genetics:
A/Gene: ovo
A/Cross-references: FlyBase:FBgn0003028

Query Match 63.3%; Score 69; DB 2; Length 1028;
Best Local Similarity 85.7%; Pred. No. 4.2;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 3 AAAAAAEEAEEAEEAEEAEEA 23
Db 497 AAAAAAAAAAAAAAAAAAAAAA 517

RESULT 8
S16356
ovo protein - fruit fly (Drosophila melanogaster)
C/Species: Drosophila melanogaster
C/Date: 13-Jan-1995 #sequence_revision 13-Jan-1995 #text_change 16-Feb-1997
C/Accession: S16356
R/Mevel-Ninio, M.; Terracol, R.; Kafatos, F.C.
EMBO J. 10, 2259-2266, 1991
A/Title: The ovo gene of Drosophila encodes a zinc finger protein required for female sex
A/Reference number: S16356; MUID:91293102; PMID:1712294
A/Accession: S16356
A/Status: preliminary
A/Molecule type: DNA
A/Residues: 1-1213 <MEV>
A/Cross-references: EMBL:X59772
C/Genetics:
A/Gene: FlyBase:ovo
A/Cross-references: FlyBase:FBgn0003028
A/Cross-references: 931/3; 1152/3

Query Match 63.3%; Score 69; DB 2; Length 1213;
Best Local Similarity 85.7%; Pred. No. 4.7;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 3 AAAAAAEEAEEAEEAEEAEEA 23
Db 860 AAAAAAAAAAAAAAAAAAAAAA 880

RESULT 9
T13748
sex comb protein - fruit fly (Drosophila melanogaster)
C/Species: Drosophila melanogaster
C/Date: 13-Aug-1999 #sequence_revision 13-Aug-1999 #text_change 17-Nov-2000
C/Accession: T13748
R/Sinclair, D.A.R.; Milne, T.A.; Hodgson, J.W.; Shellard, J.; Salinas, C.A.; Kyba, M.; R
Development 125, 1207-1216, 1998
A/Title: The additional sex combs gene of Drosophila encodes a chromatin protein that b
A/Reference number: Z17750; MUID:98146384; PMID:9477319
A/Accession: T13748
A/Status: preliminary; translated from GB/EMBL/DBJ
A/Molecule type: mRNA
A/Residues: 1-1668 <SIN>
A/Cross-references: EMBL:AJ001164; NID:93292938; PIDN:CAA04568.1; PID:93292933
C/Genetics:
A/Cross-references: FlyBase:FBgn0000142
C/Function:
A/Description: involved in repression of homeotic loci

Query Match 63.3%; Score 69; DB 2; Length 1668;

Best Local Similarity 78.3%; Pred. No. 6;
Matches 18; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

Qy 2 EAAAAAEEAEEAEEAEEAEEA 24
Db 128 KAAAAAAAAAAAAAAAAAAAAAQA 150

RESULT 10
A81882
probable dihydroilipoamide S-succinyltransferase (EC 2.3.1.61) E2 component NMA1150 [impo
C/Species: Neisseria meningitidis
C/Date: 05-May-2000 #sequence_revision 05-May-2000 #text_change 02-Feb-2001
C/Accession: A81882
R/Parthill, J.; Achtman, M.; James, K.D.; Bentley, S.D.; Churcher, C.; Klee, S.R.; Morel
Holtz, S.; Jagels, K.; Leather, S.; Moule, S.; Mungall, K.; Quail, M.A.; Rajandream,
Nature 404, 502-506, 2000
A/Title: Complete DNA sequence of a serogroup A strain of Neisseria meningitidis Z2491.
A/Reference number: A81775; MUID:20222556; PMID:10761919
A/Accession: A81882
A/Status: preliminary
A/Molecule type: DNA
A/Residues: 1-403 <PAR>
A/Cross-references: GB:AL162755; GB:AL157959; NID:97379742; PIDN:CA884412.1; PID:97379843
A/Experimental source: serogroup A, strain Z2491
C/Genetics:
A/Gene: sucB, NMA1150
C/Superfamily: dihydroilipoamide acetyltransferase; lipoyl/biotin-binding homology
C/Keywords: acyltransferase; coenzyme A

Query Match 62.4%; Score 68; DB 2; Length 403;
Best Local Similarity 75.0%; Pred. No. 2.7;
Matches 18; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

Qy 2 EAAAAAEEAEEAEEAEEAEEA 25
Db 85 EAPAAATAAEAEPAAAPAAAPAA 108

RESULT 11
A29945
neurogenesis regulatory protein - fruit fly (Drosophila melanogaster) (fragment)
N/Alternate names: single-minded gene protein
C/Species: Drosophila melanogaster
C/Date: 15-Dec-1988 #sequence_revision 15-Dec-1988 #text_change 20-Mar-1998
C/Accession: A29945
R/Crews, S.T.; Thomas, J.B.; Goodman, C.S.
Cell 52, 143-151, 1988
A/Title: The Drosophila single-minded gene encodes a nuclear protein with sequence simil
A/Reference number: A29945; MUID:88151023; PMID:3345560
A/Accession: A29945
A/Molecule type: mRNA
A/Residues: 1-655 <CRE>
A/Cross-references: GB:M19020; NID:9158464; PID:9158465
C/Genetics:
A/Gene: sim
A/Cross-references: FlyBase:FBgn0004666
C/Keywords: DNA binding; transcription regulation

Query Match 62.4%; Score 68; DB 2; Length 655;
Best Local Similarity 66.7%; Pred. No. 3.8;
Matches 16; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

Qy 2 EAAAAAEEAEEAEEAEEAEEA 25
Db 366 QAQAQAQAQAQAQAQAQAQAQA 389

RESULT 12
B53225
ecdysone-induced protein E74A - fruit fly (Drosophila virilis)
C/Species: Drosophila virilis
C/Date: 12-May-1994 #sequence_revision 12-May-1994 #text_change 21-Feb-1997

C/Accession: B53225
 R:Jones, C.W.; Dalton, M.W.; Tomkley, L.H.
 A>Title: Inter-specific comparisons of the structure and regulation of the Drosophila ecd
 A:Reference number: A53225; MUID:91200627; PMID:2016053
 A:Accession: B53225
 A>Status: preliminary; not compared with conceptual translation
 A:Molecule type: mRNA
 A:Residues: 1-873 <JON->
 A:Cross-references: GB:X59493
 C/Genetics:
 A:Gene: FlyBase: Dvir/Bip74EF
 A:Cross-references: FlyBase: FBgn0013076
 C:Superfamily: ets DNA-binding domain homology
 F:779-855/Domain: ets DNA-binding domain homology <ETS>

Query Match 62.4%; Score 68; DB 2; Length 873;
 Best Local Similarity 73.9%; Pred. No. 4.7;
 Matches 17; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 3 AAAAAAAAAAAAAAAAAAAAAA 25
 Db 501 AAAAAAAAAAATGSAAAAAAAAA 523

RESULT 13
 T33154
 polycomb protein enhancer - fruit fly (Drosophila melanogaster)
 C/Species: Drosophila melanogaster
 C/Date: 13-Aug-1999 #sequence_revision 13-Aug-1999 #text_change 17-Nov-2000
 C/Accession: T33154
 R:Stanekun, K.; Berger, J.; Ruse, C.; Sinclair, D.A.; Randazzo, F.; Brock, H.W.
 Development 125, 4055-4066, 1998
 A>Title: The enhancer of polycomb gene of Drosophila encodes a chromatin protein conserv
 A:Reference number: Z17611; MUID:98407961; PMID:9735366
 A:Accession: T33154
 A>Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: mRNA
 A:Residues: 1-2023 <STN>
 A:Cross-references: EMBL:AF079764; NID:93757889; PID:93757890; PIDN:AA64271.1
 A:Experimental source: imaginal disc
 C/Genetics:
 A:Gene: B(Pc)
 A:Cross-references: FlyBase: FBgn0000581
 A:Map position: 2

Query Match 62.4%; Score 68; DB 2; Length 2023;
 Best Local Similarity 60.0%; Pred. No. 8.7;
 Matches 18; Conservative 3; Mismatches 3; Indels 6; Gaps 1;

QY 1 CE-----AAAAAAAAAAAAAAAAAAAA 24
 Db 810 CEDQPVASTSAAAAAAAAAAAAAAAAAAS 839

RESULT 14
 JCS273
 paired type homeobox protein, NBP - human
 C/Species: Homo sapiens (man)
 C/Date: 16-Apr-1997 #sequence_revision 09-May-1997 #text_change 24-Sep-1999
 C/Accession: JCS273
 R:Yokoyama, M.; Nishi, Y.; Yoshii, J.; Okubo, K.; Matsubara, K.
 DNA Res. 3, 311-320, 1996
 A>Title: Identification and cloning of neuroblastoma-specific and nerve tissue-specific
 A:Reference number: JCS273; MUID:97191543; PMID:9039501
 A:Contents: neuroblastoma cell
 A:Accession: JCS273
 A>Status: nucleic acid sequence not shown
 A:Molecule type: mRNA
 A:Residues: 1-314 <YOK>
 A:Cross-references: DBJ:D82344; NID:91843337; PIDN:BA11555.1; PID:d1012222; PID:918413
 C:Comment: This protein is a transcriptional repressor involved in regulating gene expre
 C:Superfamily: unassigned homeobox proteins; homeobox homology

C/Keywords: DNA binding; homeobox; nucleus; transcription regulation
 F:99-155/Domain: homeobox homology <HOX>

Query Match 61.5%; Score 67; DB 2; Length 314;
 Best Local Similarity 78.3%; Pred. No. 2.8;
 Matches 18; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 3 AAAAAAAAAAAAAAAAAAAAAA 25
 Db 244 AAAAAAAAAAAAAAAAAAGGLAAA 266

RESULT 15
 T33110
 hypothetical protein C18H7.3 - Caenorhabditis elegans
 C/Species: Caenorhabditis elegans
 C/Date: 29-Oct-1999 #sequence_revision 29-Oct-1999 #text_change 21-Jan-2000
 C/Accession: T33110
 R:Tin-Wollam, A.; Fronick, W.
 submitted to the EMBL Data Library, May 1998
 A>Description: The sequence of C. elegans cosmid C18H7.
 A:Reference number: Z21284
 A:Accession: T33110
 A>Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-460 <TIN>
 A:Cross-references: EMBL:AF067607; PIDN:AACT7641.1; GSPDB:GN00022; CESP:C18H7.3
 A:Experimental source: strain Bristol N2; clone C18H7
 C/Genetics:
 A:Gene: CESP:C18H7.3
 A:Map position: 4
 A:Introns: 84/1
 C:Superfamily: Phaseolus glycine-rich cell wall protein 1.8

Query Match 61.5%; Score 67; DB 2; Length 460;
 Best Local Similarity 62.5%; Pred. No. 3.7;
 Matches 20; Conservative 0; Mismatches 4; Indels 8; Gaps 1;

QY 2 EAAAAAAAAA-----AAAAAAAAAA 25
 Db 399 EAAAAPEAAPAAEGAGGAEPAAGAAPEAAAA 430

Search completed: January 30, 2004, 00:26:21
 Job time : 11.6808 secs

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OM protein - protein search, using sw model

Run on: January 29, 2004, 23:58:27 ; Search time 5.6338 Seconds
(without alignments)
208.681 Million cell updates/sec

Title: US-09-461-684c-2
Perfect score: 109
Sequence: 1 CEAATAAEEAATAAEEAATAAEEAATAA 25

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues
Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_41:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	79	72.5	332	1	PA9A_HUMAN
2	75.5	63.3	518	1	TPM4_DROME
3	72	66.1	165	1	RS16_CAUCR
4	70	64.2	183	1	OLEC3_BRAVA
5	70	64.2	401	1	HME1_MOUSE
6	69	63.3	97	1	ANP_LIMFE
7	69	63.3	1028	1	OVO_DROME
8	69	63.3	1669	1	ASX_DROME
9	68	62.4	697	1	SIW_DROME
10	68	62.4	1073	1	HR38_DROME
11	67	61.5	314	1	PKXB_HUMAN
12	67	61.5	314	1	PKXB_MOUSE
13	67	61.5	477	1	MA2_HUMAN
14	67	61.5	606	1	HMID_DROAN
15	66	60.6	386	1	HXAD_MOUSE
16	66	60.6	388	1	HXAD_HUMAN
17	65	59.6	85	1	ANP4_PSEAM
18	65	59.6	91	1	ANPX_PSEAM
19	65	59.6	276	1	ANPX_PSEAM
20	65	59.6	280	1	SK21_HUMAN
21	65	59.6	475	1	SVX2_MOUSE
22	65	59.6	476	1	SVX2_HUMAN
23	65	59.6	644	1	BTJ_DROME
24	64	58.7	109	1	RLAI_TRYCR
25	64	58.7	392	1	HME1_HUMAN
26	64	58.7	1533	1	PUM_DROME
27	64	58.7	2038	1	FSH_DROME
28	63	57.8	179	1	RLI9_AGRIS
29	63	57.8	376	1	FXE1_HUMAN
30	63	57.8	521	1	RUN2_HUMAN
31	63	57.8	562	1	ARX_HUMAN
32	63	57.8	590	1	HMAA_DROME
33	63	57.8	607	1	RUN2_MOUSE

ALIGNMENTS

34	63	57.8	1095	1	PIPA_DROME	P13217 drosophila
35	62	56.9	364	1	NKG1_MESAU	O60554 mesocricetu
36	62	56.9	365	1	NKG1_MOUSE	O99ma9 mus musculu
37	62	56.9	365	1	NKG1_RAT	O35762 rattus norv
38	62	56.9	1355	1	SALM_DROME	P35770 drosophila
39	61	56.0	31	1	ANP3_PAGOBO	P02732 pagothenia
40	61	56.0	91	1	ANPY_PSEAM	P23699 pseudopleur
41	61	56.0	153	1	RS16_BIFLO	O86791 bifidobacte
42	61	56.0	308	1	ABE1_DROME	P38413 drosophila
43	61	56.0	376	1	FXL2_HUMAN	P58012 homo sapien
44	61	56.0	421	1	PO41_MOUSE	P17208 mus musculu
45	61	56.0	423	1	PO41_HUMAN	Q01851 homo sapien

RESULT 1
FA9A_HUMAN STANDARD; PRT; 332 AA.
ID FA9A_HUMAN
AC O81ZIU
DT 15-SEP-2003 (Rel. 42, Created)
DT 15-SEP-2003 (Rel. 42, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Protein FAM9A.
GN FAM9A.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SOURCE FROM N.A., TISSUE SPECIFICITY, AND SUBCELLULAR LOCATION.
RX MEDLINE=22202142; PubMed=12213195;
RA Martinez-Garay I., Jablonka S., Sutajova M., Steuermagel P., Gal A.,
RA Kutsche K.;
RT "A new gene family (FAM9) of low-copy repeats in Xp22.3 expressed
RT exclusively in testis: implications for recombinations in this
RT region."
RL Genomics 80:259-267(2002).
CC - SUBCELLULAR LOCATION: Nuclear; nucleolar.
CC - TISSUE SPECIFICITY: Expressed exclusively in testis.
CC - SIMILARITY: Belongs to the FAM9 family.

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CC EMBL; AF494343; AA07162.1; -
DR Genew; HGNC:18403; FAM9A.
DR Nuclear protein.
FT 180 275 GLU-RICH.
FT DOMAIN 194 214 POLY-ALA.
FT DOMAIN 252 258 POLY-GLY.
SQ SEQUENCE 332 AA; 37339 MW; 92F22EC36038229C CRC64;

Query Match 72.5%; Score 79; DB 1; Length 332;
Best Local Similarity 83.3%; Pred. No. 0.066;
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

OY 2 EAAATAAEEAATAAEEAATAAEEAATAA 25
DB 190 EAEAEAEAEAEAEAEAEAEAEAEAEAE 213

RESULT 2
TPM4_DROME STANDARD; PRT; 518 AA.
ID TPM4_DROME
AC P49455; P49456; Q24425; Q24426;
DT 01-FEB-1996 (Rel. 33, Created)

DR 01-FEB-1996 (Rel. 33, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Tropomyosin 1, isoforms 33/34 (Tropomyosin II).
 GN Tm1 OR Tm1.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 CC Neoptera; Endopterygota; Diptera; Brachyera; Muscomorpha;
 CC Ephydriidae; Drosophilidae; Drosophila.
 NC NCB1_Taxid=7227;
 RN (1)
 RP SEQUENCE FROM N.A. (ISOFORMS 33 AND 34).
 RC TISSUE=Embryo, and Pupae;
 RX MEDLINE=89127197; PubMed=2851721;
 RA Hanke P.D., Scott R.V.;
 RT "The Drosophila melanogaster tropomyosin II gene produces multiple
 RT proteins by use of alternative tissue-specific promoters and
 RT alternative splicing."
 RT Mol. Cell. Biol. 8:3591-3602(1988).
 RL (2)
 RP SEQUENCE FROM N.A. (ISOFORMS 33 AND 34).
 RC STRAIN=Oregon-R; TISSUE=Pupae;
 RX MEDLINE=87064486; PubMed=3097506;
 RA Karlik C.C., Fyrborg E.A.;
 RT "Two Drosophila melanogaster tropomyosin genes: structural and
 RT functional aspects."
 RT Mol. Cell. Biol. 6:1965-1973 (1986).
 RL (1)
 RP ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=5;
 CC Comment=Additional isoforms seem to exist;
 CC Name=33; Synonym=9C;
 CC IsoId=P06754-1; Sequence=Displayed;
 CC Name=Muscle; Synonym=9D;
 CC IsoId=P06754-1; Sequence=External;
 CC Name=Non-muscle; Synonym=Cytoskeletal;
 CC IsoId=P06754-2; Sequence=External;
 CC Name=9A;
 CC IsoId=P06754-3; Sequence=External;
 CC Name=34; Synonym=9B;
 CC IsoId=P06754-2; Sequence=VSP_006623, VSP_006624, VSP_006625;
 CC -1- TISSUE SPECIFICITY: Both isoforms are only expressed in indirect
 CC flight muscles.
 CC -1- DEVELOPMENTAL STAGE: Both isoforms are expressed during pupal and
 CC adult stages.
 CC -1- DOMAIN: THE MOLECULE IS IN A COILED COIL STRUCTURE. THE SEQUENCE
 CC EXHIBITS A PROMINENT SEVEN-RESIDUES PERIODICITY.
 CC -1- SIMILARITY: BELONGS TO THE TROPOMYOSIN FAMILY.
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 CC -----
 DR EMBL: X76208; CAA53800.1; -;
 DR EMBL: X76208; CAA53801.1; -;
 DR EMBL: K02620; AAA28967.1; ALT_SEQ.
 DR EMBL: K02620; AAA28967.1; JOINED.
 DR EMBL: L00355; AAA28967.1; JOINED.
 DR EMBL: L00355; AAA28967.1; JOINED.
 DR EMBL: L00357; AAA28967.1; JOINED.
 DR EMBL: L00357; AAA28967.1; JOINED.
 DR EMBL: L00359; AAA28967.1; JOINED.
 DR EMBL: L00359; AAA28967.1; JOINED.
 DR EMBL: L00360; AAA28967.1; JOINED.
 DR EMBL: L00362; AAA28967.1; JOINED.
 DR EMBL: L00362; AAA28967.1; JOINED.
 DR EMBL: M12840; AAA28967.1; JOINED.
 DR EMBL: M12840; AAA28967.1; JOINED.
 DR EMBL: K02621; AAA28968.1; -;
 DR EMBL: M12840; AAA28968.1; JOINED.
 DR EMBL: L00355; AAA28968.1; JOINED.
 DR EMBL: L00355; AAA28968.1; JOINED.
 DR EMBL: L00356; AAA28968.1; JOINED.
 DR EMBL: L00357; AAA28968.1; JOINED.
 DR EMBL: L00358; AAA28968.1; JOINED.
 DR EMBL: L00359; AAA28968.1; JOINED.
 DR EMBL: L00359; AAA28968.1; JOINED.

DR EMBL: L00360; AAA28968.1; JOINED.
 DR EMBL: L00362; AAA28968.1; JOINED.
 DR FlyBase; FBgn003721; Tm1.
 DR InterPro; IPR00533; Tropomyosin.
 DR Pfam; PF00261; Tropomyosin.1.
 DR PRINTS; PR00194; TROPOMYOSIN.
 DR PROSITE; PS00326; TROPOMYOSIN; 1.
 KW Muscle protein; Cytoskeleton; Actin-binding; coiled coil;
 KM Alternative splicing; Multigene family.
 FT DOMAIN
 FT 14
 FT 267
 FT 287
 FT 293
 FT VARSPLIC
 FT 300
 FT 367
 FT VARSPLIC
 FT 391
 FT 518
 FT VARSPLIC
 FT 518
 FT 106
 FT 114
 FT 119
 FT 183
 FT 199
 FT 503
 FT 503
 FT 5458 MW; 1530872CF9DB6EA CXC64;
 SQ SEQUENCE
 Query Match
 Best Local Similarity 69.3%; Score 75.5; DB 1; Length 518;
 Matches 21; Conservative 0; Mismatches 2; Indels 1; Gaps 1;
 DB 2 EMBL:AAA28968.1-AAA28968.1 25
 DB 314 EMBL:AAA28968.1-AAA28968.1 336
 RESULT 3
 RS16 CAUCR STANDARD; PRT; 165 AA.
 AC P58122;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE 30S ribosomal protein S16.
 GN RPS16 OR CC3652.
 OS Caulobacter crescentus.
 OC Bacteria; Proteobacteria; Alphaproteobacteria; Caulobacteriales;
 CC Caulobacteriaceae; Caulobacter.
 NC NCB1_Taxid=155892;
 RN (1)
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC 19089 / CB15;
 RX MEDLINE=21173698; PubMed=11259647;
 RA Nielsen W.C., Feidbylum T.V., Laub M.T., Paulsen I.T., Nelson K.E.,
 RA Eriksen U., Heidelberg J.F., Alley M.R.K., Ohta N., Maddock J.R.,
 RA Potocka I., Nelson W.C., Newton A., Stephens C., Phadke N.D., Ely B.,
 RA DeBoy R.T., Dodson R.J., Durkin A.S., Gwinn M.L., Haft D.H.,
 RA Kolonay J.F., Smit J., Craven M.B., Khouli H., Shetty K.,
 RA Uitterlind T., Tran K., Wolf A., Vamathevan J., Ermolaeva M., White O.,
 RA Salzberg S.L., Venter J.C., Shapiro L., Fraser C.M.;
 RT Complete genome sequence of Caulobacter crescentus."
 RT Proc. Natl. Acad. Sci. U.S.A. 98:4136-4141(2001).
 CC -1- SIMILARITY: BELONGS TO THE S16P FAMILY OF RIBOSOMAL PROTEINS.
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DR EMBL; AE006023; AKK25614.1; --
 DR PIR; B87702; B87702.
 DR HSSP; P80379; 1EMW.
 DR TIGR; CC3652; --
 DR HAMAP; MF_00385; -- 1.
 DR InterPro; IPR000307; Ribosomal_S16.
 DR Pfam; PF00886; Ribosomal_S16; 1.
 DR ProDom; PD003791; Ribosomal_S16; 1.
 DR TIGRFAMs; TIGR00002; S16; 1.
 DR PROSITE; PS00732; RIBOSOMAL_S16; 1.
 DR Ribosomal protein, Complete proteome.
 KW Ribosomal protein, Complete proteome.
 SQ SEQUENCE 165 AA; 17605 MW; ED46FC2798C5BE1C CRC64;

Query Match 66.1%; Score 72; DB 1; Length 165;
 Best Local Similarity 75.0%; Pred. No. 0.19;
 Matches 18; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

OY 2 EAAAAAAAAAAAAAAAAAAAAA 25
 DB 115 QAEADKAAAEKXKAAAAAAAAA 138

RESULT 4
 OLEC_BRANA STANDARD; PRT; 183 AA.
 AC P29526;
 DT 01-APR-1993 (Rel. 25, Created)
 DT 01-APR-1993 (Rel. 25, Last sequence update)
 DT 15-JUL-1998 (Rel. 36, Last annotation update)
 DE Oleosin C98 (Fragment).
 GN C98
 OS Brassica napus (Rape).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
 OC eurosids II; Brassicales; Brassicaceae; Brassica.
 OX NCBI_TaxID=3708;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=anther;
 RX MEDLINE=93386188; PubMed=8374615;
 RA Roberts M.R., Hodge R., Rose J.H.E., Sorensen A., Murphy D.J.,
 RA Draper J., Scott R.;
 RT "Characterization of a new class of oleosins suggests a male
 gametocyte-specific lipid storage pathway".
 RL Plant J. 3:629-636(1993).
 RT FUNCTION: MAY HAVE A STRUCTURAL ROLE TO STABILIZE THE LIPID BODY
 DURING DESEICATION OF THE SEED BY PREVENTING COALESCENCE OF THE
 OIL. PROBABLY INTERACTS WITH BOTH LIPID AND PHOSPHOLIPID MOLECULES
 OF LIPID BODIES. MAY ALSO PROVIDE RECOGNITION SIGNALS FOR SPECIFIC
 LIPASE ANCHORAGE IN LIPOLYSIS DURING SEEDLING GROWTH.
 CC MONOLAYER LIPID/WATER INTERFACE.
 CC SUBCELLULAR LOCATION: SURFACE OF OIL BODIES. OLEOSINS EXIST AT A
 -1- TISSUE SPECIFICITY: SPECIFIC TO THE MALE GAMETOPHYTE.
 CC -1- SIMILARITY: BELONGS TO THE OLEOSIN FAMILY.
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DR EMBL; X67142; CAA47623.1; --
 DR PIR; S24960; S24960.
 DR InterPro; IPR000136; Oleosin.

DR Pfam; PF01277; Oleosin; 1.
 DR PROSITE; PS00811; OLEOSINS; 1.
 KW Seed; Oil body; Multigene family.
 FT NON_TER 1
 FT DOMAIN 1
 FT DOMAIN 23
 FT DOMAIN 24
 FT DOMAIN 95
 SQ SEQUENCE 183 AA; 18149 MW; 198A5D3B6DF3045A CRC64;

Query Match 64.2%; Score 70; DB 1; Length 183;
 Best Local Similarity 78.3%; Pred. No. 0.33;
 Matches 18; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

OY 3 AAAAAAAAAAAAAAAAAAAAAA 25
 DB 151 AAPAAPAPAPAPAPAPAPAPAA 173

RESULT 5
 HME1_MOUSE STANDARD; PRT; 401 AA.
 ID HME1_MOUSE
 AC P09065;
 DT 01-NOV-1998 (Rel. 09, Created)
 DT 01-FEB-1994 (Rel. 28, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Homeobox protein engrailed-1 (Mo-Eu-1).
 GN EN1 OR EN-1.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=93185339; PubMed=1363401;
 RA Logan C., Hanks M.C., Noble-Topham S., Nallathattan D.,
 RA Provart N.J., Joyner A.L.;
 RT "Cloning and sequence comparison of the mouse, human, and chicken
 engrailed genes reveal potential functional domains and regulatory
 regions".
 RL Dev. Genet. 13:345-358(1992).
 RN [2]
 RP SEQUENCE OF 278-401 FROM N.A.
 RX MEDLINE=88112776; PubMed=2892757;
 RA Joyner A.L., Martin G.R.;
 RT "En-1 and En-2, two mouse genes with sequence homology to the
 Drosophila engrailed gene: expression during embryogenesis".
 RL Genes Dev. 1:29-38(1987).
 RN [3]
 RP SEQUENCE OF 298-401 FROM N.A.
 RX MEDLINE=86079501; PubMed=2416459;
 RA Joyner A.L., Kornberg T., Coleman K.G., Cox D.R., Martin G.R.;
 RT "Expression during embryogenesis of a mouse gene with sequence
 homology to the Drosophila engrailed gene".
 RL Cell 43:29-37(1985).
 RN [4]
 RP SEQUENCE OF 321-380 FROM N.A.
 RX MEDLINE=91095509; PubMed=1980115;
 RA Holland P.W.H., Williams N.A.;
 RT "Conservation of engrailed-like homeobox sequences during vertebrate
 evolution".
 RL FEBS Lett. 277:250-252(1990).
 CC -1- SUBCELLULAR LOCATION: Nuclear.
 CC -1- SIMILARITY: BELONGS TO THE ENGRAILED HOMEOBOX FAMILY.
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DR EMBL; L12703; AAA03660.2; --

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DR EMBL: Y00201; CAA68361.1; -.
DR PIR: A48423; A48423.
DR HSSP: P02836; 3HSD.
DR TRANSFAC: T02016; -.
DR MGD: MGI:95389; Enl.
DR InterPro: IPR000747; Engrailed.
DR InterPro: IPR001356; Homeobox.
DR InterPro: IPR000047; HTH_Lambrprepress.
DR Pfam: PF00046; homeobox_1.
DR PRINTS: PR00026; ENGRAILED.
DR PRINTS; PR00024; HOMEBOX.
DR PRINTS; PR00031; HTHREPRESSR.
DR ProDom: PD000010; Homeobox; 1.
DR SMART: SM00389; HOX; 1.
DR PROSITE: PS00027; HOMEBOX_1; 1.
DR PROSITE; PS0071; HOMEBOX_2; 1.
DR PROSITE; PS00033; ENGRAILED; 1.
DR Homeobox; DNA-binding; Developmental protein; Nuclear protein.
FT DOMAIN 52 87 PRO-RICH.
FT DOMAIN 73 87 POLY-PRO.
FT DOMAIN 207 228 POLY-ALA.
FT DNA_BIND 312 371 HOMEBOX.
SQ SEQUENCE 401 AA; 40950 MW; 1F90210950152PAB CRC64;

Query Match 64.2%; Score 70; DB 1; Length 401;
Best Local Similarity 78.3%; Pred No. 0.62;
Matches 18; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

OY 3 EMBL: Y00201; CAA68361.1; -.
DB 207 AAAAAAAAAAAAAAAAAAAAAA 229

RESULT 6
ANP_LIMFE
ID ANP_LIMFE STANDARD; PRT; 97 AA.
AC P09031;
DT 01-NOV-1988 (Rel. 09, Last sequence update)
DT 01-NOV-1988 (Rel. 09, Last sequence update)
DT 01-AUG-1990 (Rel. 15, Last annotation update)
DE Antifreeze protein precursor (AFP).
OS Limanda ferruginea (Yellowtail flounder).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei;
OC Acanthomorpha; Acanthopterygii; Perciformes;
OC Pleuronectidae; Pleuronectidae; Limanda.
OC NCB1_Taxid=8258;
[1]
SEQUENCE FROM N.A.
RX MEDLINE=88029483; PubMed=3665937;
RA Scott G.K.; Davies P.L.; Shears M.A.; Fletcher G.L.;
RT "Structural variations in the alanine-rich antifreeze proteins of the
RT pleuronectinae."
RL Eur. J. Biochem. 168:629-633(1987).
CC -1- FUNCTION: ANTIFREEZE PROTEINS LOWER THE BLOOD FREEZING POINT.
CC -1- SIMILARITY: BELONGS TO THE TYPE-I AFP FAMILY. TYPE 1 AFP ARE
CC ALANINE-RICH, AMPHIPHILIC AND ALPHA-HELICAL.
CC
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EMBL: X06356; CAA29655.1; -.
DR PIR: S02376; S02376.
DR InterPro: IPR000104; Antifreeze_1.
DR PRINTS; PR00308; ANTIFREEZE1.
DR Antifreeze protein; Repeat; Signal.
FT SIGNAL 1 23
FT PROPEP 24 48 REMOVED BY A DIPEPTIDYL-PEPTIDASE

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FT CHAIN 49 97 (PROBABLE).
FT SEQUENCE 97 AA; 8865 MW; 62AD582DF8E45B6 CRC64;
SQ SEQUENCE 97 AA; 8865 MW; 62AD582DF8E45B6 CRC64;

Query Match 63.3%; Score 69; DB 1; Length 97;
Best Local Similarity 70.8%; Pred. No. 0.26;
Matches 17; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

OY 2 EMBL: Y00201; CAA68361.1; -.
DB 53 DAAAAAAAAATAAAKAAADTAAA 76

RESULT 7
OVO_DROME
ID OVO_DROME STANDARD; PRT; 1028 AA.
AC P51521; Q9XZU4;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Ovo protein (Shaven baby protein).
DE OVO OR SVB.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydriidae; Drosophilidae; Drosophila.
OC NCB1_Taxid=7227;
[1]
SEQUENCE FROM N.A.
RX MEDLINE=95021209; PubMed=7935398;
RA Garfinkel M.D.; Wang J.; Liang Y.; Mahowald A.P.;
RT "Multiple products from the shavenbaby-ovo gene region of Drosophila
RT melanogaster: relationship to genetic complexity."
RL Mol. Cell. Biol. 14:6809-6818(1994).
[2]
SEQUENCE FROM N.A.
RX STRAIN-Oregon-R;
RA MEDLINE=91293102; PubMed=1712294;
RA Wevel-Minto M.T.M.; Terracol R.; Kafatos F.C.;
RT "The ovo gene of Drosophila encodes a zinc finger protein required
RT for female germ line development."
RL EMBL J. 10:3259-3266(1991).
CC -1- FUNCTION: REQUIRED FOR SURVIVAL AND DIFFERENTIATION OF FEMALE GERM
CC LINE CELLS. PLAYS A ROLE IN GERM LINE SEX DETERMINATION.
CC -1- SUBCELLULAR LOCATION: Nuclear (potential).
CC ACCUMULATES IN NURSE CELLS DURING OOGENESIS. STORED IN THE EGG,
CC BUT IS RAPIDLY LOST IN THE EMBRYOS EXCEPT FOR ITS CONTINUED
CC PRESENCE IN THE GERM LINE PRECURSOR POLE CELLS.
CC -1- SIMILARITY: Contains 4 C2H2-type zinc fingers.
CC
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EMBL: U11383; AAB60216.1; -.
DR EMBL: X59772; CAB36921.1; ALT_SEQ.
DR PIR: A56038; A56038.
DR HSSP: P07248; ZADR.
DR TRANSFAC: T00669; -.
DR FlyBase: FBgn003028; ovo.
DR InterPro: IPR007087; Znf_C2H2.
DR Pfam: PF00096; zf-C2H2_3.
DR SMART; SM00355; ZNF_C2H2; 4.
DR PROSITE; PS00028; ZINC_FINGER_C2H2_1; 3.
DR PROSITE; PS00157; ZINC_FINGER_C2H2_2; 3.
DR Zinc-finger; Metal-binding; DNA-binding; Repeat; Nuclear protein;
KW Transcription regulation.

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FT DOMAIN 62 66 POLY-ALA.
FT DOMAIN 72 77 POLY-GLY.
FT DOMAIN 80 85 POLY-GLY.
FT DOMAIN 98 108 POLY-GLY.
FT DOMAIN 144 152 POLY-HIS.
FT DOMAIN 153 159 POLY-ASN.
FT DOMAIN 336 339 POLY-GLN.
FT DOMAIN 347 353 POLY-GLN.
FT DOMAIN 357 361 POLY-GLN.
FT DOMAIN 410 414 POLY-GLN.
FT DOMAIN 418 422 POLY-GLN.
FT DOMAIN 426 432 POLY-GLN.
FT DOMAIN 445 453 POLY-GLN.
FT DOMAIN 456 459 POLY-GLN.
FT DOMAIN 466 474 POLY-GLN.
FT DOMAIN 497 517 POLY-ALA.
FT DOMAIN 524 529 POLY-SER.
FT DOMAIN 539 558 POLY-ALA.
FT DOMAIN 639 651 POLY-ALA.
FT DOMAIN 717 725 POLY-ALA.
FT DOMAIN 797 802 POLY-GLN.
FT DOMAIN 820 823 POLY-GLN.
FT DOMAIN 826 832 POLY-GLN.
FT ZN FING 874 896 C2H2-TYPE 1.
FT ZN FING 902 924 C2H2-TYPE 2.
FT ZN FING 930 953 C2H2-TYPE 3.
FT ZN FING 969 992 C2H2-TYPE 4.
FT ZN FING 969 992 A -> R (IN REF. 2).
FT ZN FING 969 992 A -> R (IN REF. 2).
SQ SEQUENCE 1028 AA; 110620 MW; D7068B2BC0F677 CRC64;

Query Match
Best Local Similarity 63.3%; Score 69; DB 1; Length 1028;
Matches 18; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 3 AAAAAAAAAAAAAAAAAAAAAA 23
DB 497 AAAAAAAAAAAAAAAAAAAAAA 517

RESULT 8
ASX DROME STANDARD; PRT; 1669 AA.
AC 09V727; 076930;
DT 15-SEP-2003 (Rel. 42, Created)
DT 15-SEP-2003 (Rel. 42, Last sequence update)
DE Polycomb protein Asx (Additional sex combs).
GN ASX OR CG8787.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydriidae; Drosophilidae; Drosophila.
OC NCBI TaxID=7227;
RP SEQUENCE FROM N.A., FUNCTION, SUBCELLULAR LOCATION, AND DEVELOPMENTAL
RP STAGE.
RC TISSUE=Imaginal disks;
RC MEDLINE=9814684; PubMed=9477319;
RA Simlaid D.A.R., Milne T.A., Hodgson J.W., Shellard J., Salinas C.A.,
RA Kyba M., Randazzo F., Brock H.W.;
RA "The Additional sex combs gene of Drosophila encodes a chromatin
RT protein that binds to shared and unique Polycomb group sites on
RT polytene chromosomes.";
RT Development 125:1207-1216 (1998).
RL [2]
RP SEQUENCE FROM N.A.
RC STRAIN=Berkeley;
RC MEDLINE=20196006; PubMed=10731132;
RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
RA Sutton G.G., Mortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
RA Brandon R.C., Rogers Y.-H.C., Blazej R.G., Champe M., Pfeiffer B.D.,

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RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
RA Beeson K.Y., Benos P.V., Bernier B.P., Bhandari D., Bolshakov S.,
RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brotlier P.,
RA Burris K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
RA de Pablo S., DeCher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
RA Durbin K.D., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
RA Foster C., Gabriellian A.E., Garg N.S., Gelbart W.M., Glasser K.,
RA Glodok A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibeagwam C.,
RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
RA Lasok P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
RA Merkulov G., Milhina N.V., Mobarry C., Morris J., Moshrefi A.,
RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
RA Nelson D.R., Nelson K.A., Nixon K., Nusken D.R., Paclet J.M.,
RA Palazzolo M., Peltman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
RA Spier B., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weissbach J.,
RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
RT "The genome sequence of Drosophila melanogaster.";
RL Science 287:2185-2195 (2000).
RN [3]
RP INTERACTION WITH TAN.
RX MEDLINE=21290825; PubMed=11397012;
RA Dietrich B.H., Moore J., Kyba M., dosSantos G., McClokey F.,
RA Milne T.A., Brock H.W., Krause H.M.;
RT "Tantalus", a novel ASX-interacting protein with tissue-specific
RT functions.";
RL Dev. Biol. 234:441-453 (2001).
CC -1- FUNCTION: Atypical Polycomb group protein, which may be involved
CC in both Polycomb group (PcG) and trithorax group (trxG) complexes.
CC PcG and trxG proteins act by forming multiprotein complexes, which
CC are respectively required to maintain the transcriptionally
CC repressive and transcriptionally active state of homeotic genes
CC throughout development. PcG and trxG protein complexes are not
CC required to initiate repression and activation, but to maintain it
CC during later stages of development. Both complexes probably act
CC via methylation of histones, rendering chromatin heritably changed
CC in its expressibility.
CC -1- SUBUNIT: Interacts with Tan.
CC -1- SUBCELLULAR LOCATION: Nuclear; associated with chromatin.
CC Colocalizes with many PcG sites on polytene chromosomes. It also
CC associates with many unique sites on polytene chromosomes.
CC -1- TISSUE SPECIFICITY: Highly expressed in nurse cells and deposited
CC in oocytes late in oogenesis. Ubiquitous in early embryos. Late
CC embryos show higher levels in CNS and neuroectoderm.
CC -1- DEVELOPMENTAL STAGE: Expressed both maternally and zygotically.
CC Early embryos have high levels of expression, this drops off and
CC zygotic expression begins at 3-6 hour embryos. Expression levels
CC are low in larvae and medium in pupae and adults.
CC -1- DOMAIN: Contains two Leu-Xaa-Leu-Leu (LXLL) motifs, which may
CC be required for an association with nuclear receptors (By
CC similarity).
CC -1- SIMILARITY: Belongs to the Asx family.
CC -1- SIMILARITY: Contains 1 PHD-type zinc finger.
CC -1- CAUTION: Ref.1 sequence differs from that shown due to
CC frameshifts in positions 608 and 719.
CC -----
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CC -----
 DR EMBL: AJ001164; CA04568.1; ALT_FRAME.
 DR EMBL: AE003814; AAF58239.1; -.
 DR FLYBASE: Fgn0000142; Asx.
 DR PROSITE: PS01359; ZF_PHD_1; FALSE_NEG.
 DR PROSITE: PS50016; ZF_PHD_2; FALSE_NEG.
 KM Transcription regulation: Repressor; Nuclear protein; zinc;
 KW Metal-binding; Zinc-finger; Repeat; Developmental protein.
 FT ZN_FING 1632 1669
 FT DOMAIN 8 12
 FT DOMAIN 122 126 POLY-GLN.
 FT DOMAIN 129 152 POLY-GLN.
 FT DOMAIN 638 715 ALA-RICH.
 FT DOMAIN 747 751 SER-RICH.
 FT DOMAIN 862 1202 POLY-GLN.
 FT DOMAIN 1287 1290 GLN-RICH.
 FT DOMAIN 1520 1524 POLY-THR.
 FT DOMAIN 1527 1536 POLY-HIS.
 FT SITE 224 228 LKLL MOTIF 1.
 FT SITE 244 248 LKLL MOTIF 2.
 FT CONFLICT 14 15 SQ -> CE (IN REF. 1).
 FT CONFLICT 187 187 K -> N (IN REF. 1).
 FT CONFLICT 1253 1253 S -> T (IN REF. 1).
 FT CONFLICT 1520 1520 MISSING (IN REF. 1).
 SQ SEQUENCE 1669 AA; F65D67398D6D321 CRC64;

Query Match 63.3%; Score 69; DB 1; Length 1669;
 Best Local Similarity 78.3%; Pred. No. 2.4;
 Matches 18; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 2 EAAAAAAAAAAAAAAAAAAAAA 24
 Db 128 KAAAAAAAAAAAAAAAAAAAAA 150

RESULT 9
 SIM_DROME STANDARD; PRT; 697 AA.
 ID SIM_DROME
 AC P05709; O96521; Q8MQ17; Q9VEZ3;
 DT 01-NOV-1988 (Rel. 09, Created)
 DT 15-SEP-2003 (Rel. 42, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Single-minded protein.
 GN SIM OR CG7771.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 CC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 CC Ephydroidea; Drosophilidae; Drosophila.
 OK NCBI_TaxID=7227;
 (1)
 RP SEQUENCE FROM N.A.
 RP MEDLINE=99054545; PubMed=9840810;
 RA Kasai Y., Stahl S., Crews S.;
 RT "Specification of the Drosophila CNS midline cell lineage: direct
 RT control of single-minded transcription by dorsal/ventral patterning
 RT genes";
 RL Gene Expr. 7:171-189(1998).
 (2)
 RP SEQUENCE FROM N.A.
 RP STRAIN=Berkley;
 MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amaralides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galle R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Suton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.-H.C., Blaise R.G., Chame M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Makos G.L.G.,
 RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
 RA Ballew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,

RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Broctier P.,
 RA Burris K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Fabios B., Delcher A., Deng Z., Nays A.D., Dew I., Dietz S.M.,
 RA Dodeon K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durkin K.J., Evangelista C.C., Ferraz C., Ferriera S., Fleischmann W.,
 RA Foster C., Gabrieli A.E., Gary N.S., Gelbart W.M., Glaeser K.,
 RA Glodok A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.U., Wei M.-H., Ibegwan C.,
 RA Jallali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Lascko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Noy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Pacle J.M.,
 RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Svirskas R., Tector R., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Wassarman D.A., Weinstein G.M., Weissensbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.,
 RT "The genome sequence of Drosophila melanogaster.";
 RL Science 287:2185-2195(2000).
 (3)
 RP REVISIONS.
 RP STRAIN=Berkley;
 MEDLINE=22426069; PubMed=12537572;
 RC STRAIN=Berkley;
 RX Mista S., Crosby M.A., Mungall C.J., Matthews B.B., Campbell K.S.,
 RA Hradecky P., Huang Y., Kaminker J.S., Milburn G.H., Prochuk S.E.,
 RA Smith C.D., Tupy J.L., Whitfield E.J., Bayraktaroglu L., Berman B.P.,
 RA Bettencourt B.R., Celniker S.E., de Grey A.D.N.J., Drysdale R.A.,
 RA Harris N.L., Richter J., Russo S., Schroeder A.J., Shu S.O.,
 RA Stapleton M., Yamada C., Ashburner M., Gelbart W.M., Rubin G.M.,
 RA Lewis S.E.;
 RT "Annotation of the Drosophila melanogaster euchromatic genome: a
 RT systematic review";
 RL Genome Biol. 3:RESEARCH0083.1-RESEARCH0083.22(2002).
 (4)
 RP SEQUENCE FROM N.A.
 RP STRAIN=Berkley; TISSUE=Embryo;
 MEDLINE=22426066; PubMed=12537569;
 RX Stapleton M., Carlson J.W., Brokstein P., Yu C., Chame M.,
 RA George R.A., Guarin H., Kronmiller B., Pacle J.M., Park S., Wan K.H.,
 RA Rubin G.M., Celniker S.E.;
 RT "A Drosophila full-length cDNA resource";
 RL Genome Biol. 3:RESEARCH0080.1-RESEARCH0080.8(2002).
 (5)
 RP SEQUENCE OF 25-42 FROM N.A., AND SIMILARITY TO HLH PROTEINS.
 RP MEDLINE=92103681; PubMed=1760843;
 RA Nabu J.R., Lewis J.O., Wharton K.A., Jr., Crews S.T.;
 RT "The Drosophila single-minded gene encodes a helix-loop-helix protein
 RT that acts as a master regulator of CNS midline development";
 RL Cell 67:1157-1167(1991).
 (6)
 RP SEQUENCE OF 43-697 FROM N.A.
 RP MEDLINE=88151023; PubMed=3345560;
 RA Crews S.T., Thomas J.B., Goodman C.S.;
 RT "The Drosophila single-minded gene encodes a nuclear protein with
 RT sequence similarity to the per gene product";
 RL Cell 52:143-151(1988).
 CC -1- FUNCTION: TRANSCRIPTION FACTOR THAT FUNCTIONS AS A MASTER
 CC DEVELOPMENTAL REGULATOR OF THE CNS MIDLINE LINEAGE. MUTATIONS IN
 CC THE SIM GENE RESULTS IN THE LOSS OF THE PRECURSOR CELLS GIVING
 CC RISE TO MIDLINE CELLS OF THE EMBRYONIC CENTRAL NERVOUS SYSTEM.
 CC -1- SUBUNIT: Efficient DNA binding requires dimerization with another
 CC HLH protein.

CC -1- SUBCELLULAR LOCATION: Nuclear (Potential).
 CC -1- POLYMORPHISM: Berkeley strain has 11 A-A-Q repeats.
 CC -1- SIMILARITY: BELONGS TO THE BASIC HELIX-LOOP-HELIX (BHLH) FAMILY OF
 CC TRANSCRIPTION FACTORS.
 CC -1- SIMILARITY: Contains 2 PAS (PER-ARNT-SIM) dimerization domains.
 CC -1- SIMILARITY: Contains 1 PAS-associated C-terminal (PAC) domain.
 CC -1- CAUTION: Ref.1 sequence differs from that shown due to erroneous
 CC gene model prediction.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL, AF071934; AAC64519.1; ALT_SBO.
 CC EMBL, AB003698; AAF54902.3; -.
 CC EMBL, AY129457; AAM76199.1; -.
 CC EMBL, M19020; AAM28900.1; -.
 CC PIR, A29945; A29945.
 CC TRANSFAC, T00750; -.
 CC FLYBase, FBgn0004666; sim.
 CC DR GO, GO:0005634; C:nucleus, IEP.
 CC DR GO, GO:0003702; F:RNA polymerase II transcription factor acti., NMS.
 CC DR GO, GO:0006355; P:regulation of transcription, DNA-dependent, NMS.
 CC DR GO, GO:0007418; P:ventral midline development, IMP.
 CC DR InterPro, IPR001092; HLH_baslc.
 CC DR InterPro, IPR001067; Nuc_translocat.
 CC DR InterPro, IPR001610; PAC.
 CC DR InterPro, IPR000014; PAS_domain.
 CC DR Pfam, PF00010; HLH; 1.
 CC DR Pfam, PF00785; PAC; 1.
 CC DR Pfam, PF00989; PAS; 1.
 CC DR PRINTS, PR00785; NCTRNLOCATR.
 CC DR SMART, SMO0353; HLH; 1.
 CC DR SMART, SMO0086; PAC; 1.
 CC DR SMART, SMO0091; PAS; 2.
 CC DR TIGRFAMs, TIGR00229; sensory_box; 2.
 CC DR PROSITE, PS00038; HLH_1; 1.
 CC DR PROSITE, PS00888; HLH_2; 1.
 CC DR PROSITE, PS0112; PAS; 2.
 CC KW Developmental protein; Neurogenesis; Nuclear protein; Repeat;
 CC Transcription regulation; DNA-binding.
 CC FT DNA_BIND 21 37
 CC FT DOMAIN 38 78 BASIC DOMAIN.
 CC FT DOMAIN 100 172 HELIX-LOOP-HELIX MOTIF (BY SIMILARITY).
 CC FT DOMAIN 266 336 PAS 1.
 CC FT DOMAIN 406 446 PAS 2.
 CC FT DOMAIN 553 672 14 X 3 AA REPEATS OF A-A-Q (APPROXIMATE).
 CC FT DOMAIN 673 693 SER-RICH.
 CC FT DOMAIN 693 693 GLN/HIS-RICH.
 CC FT VARIANT 406 414 MISSING (IN STRAIN BERKELEY).
 CC FT CONFLICT 151 151 I -> Y (IN REF. 1).
 CC SQ SEQUENCE 697 AA; 76475 MW; 58841AA4A17101AD CRC64;

Query Match 62.4% Score 68 DB 1; Length 697;
 Best Local Similarity 66.7% Pred. No. 1.5;
 Matches 16; Conservative 4; Mismatches 4; Indels 0; Gaps 0;
 Oy 2 EAAAAAAAAAAAAAAAAAAAAA 25
 Db 408 QAAQAAQAAQAAQAAQAAQAA 431

RESULT 10
 ID HR38 DROME STANDARD; PRT; 1073 AA.
 AC P49869; O18383; Q9V1K4;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Probable nuclear hormone receptor HR38 (DHR38).

GN HR38 OR NR44 OR CG1864.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A. (ISOFORM SHORT).
 RC TISSUE=Larva;
 RX MEDLINE=95372400; PubMed=7644522;
 RA Sutherland J.D., Kozlova T., Tzertzinis G., Kafatos F.C.;
 RT "Drosophila hormone receptor 38: a second partner for Drosophila USP
 RT suggests an unexpected role for nuclear receptors of the nerve growth
 RT factor-induced protein B type.";
 RT Proc. Natl. Acad. Sci. U.S.A. 92:7966-7970(1995).
 RN [2]
 RP SEQUENCE FROM N.A. (ISOFORM SHORT), AND TISSUE SPECIFICITY.
 RX MEDLINE=98370123; PubMed=9704500;
 RA Komonyi O., Mink M., Csiba J., Maroy P.;
 RT "Genomic organization of DHR38 gene in Drosophila: presence of
 RT Alu-like repeat in a translated exon and expression during embryonic
 RT development.";
 RL Arch. Insect Biochem. Physiol. 38:185-192(1998).
 RN [3]
 RP SEQUENCE FROM N.A. (ISOFORM LONG), AND CHARACTERIZATION.
 RC TISSUE=Larva;
 RX MEDLINE=98315108; PubMed=9649534;
 RA Kozlova T., Pokholkova G.V., Tzertzinis G., Sutherland J.D.,
 RA Zhmulev I.F., Kafatos F.C.;
 RT "Drosophila hormone receptor 38 functions in metamorphosis: a role in
 RT adult cuticle formation.";
 RL Genetics 149:1465-1475(1998).
 RN [4]
 RP SEQUENCE FROM N.A. (ISOFORM LONG).
 RC STRAIN=Berkeley;
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celiker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hopkins R.A., Galie R.E.,
 RA Sutton R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Geaton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.-H.C., Blazej R.G., Champe M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
 RA Abril J.F., Aghayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
 RA Ballew R.M., Baxevanis A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brotter P.,
 RA Butts K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cavley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablo J., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunn P.,
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
 RA Foster C., Gabriellian A.E., Garg N.S., Gelbart W.M., Glasner K.,
 RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,
 RA Jalali M., Kalush F., Karpen G.H., Ke Z., Kemison J.A., Ketchum K.A.,
 RA Kimmel B.B., Kodra C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Laako P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nuskern D.R., Pacle J.M.,
 RA Palazolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Klamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier B., Spradling A.C., Stapleton M., Strong R., Sun B.,
 RA Svrlakas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weissbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of Drosophila melanogaster."

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RL [5] Sequence: 287:2185-2195 (2000).
RN
RP SEQUENCE OF 528-1073 FROM N.A.
RC STRAIN=Canon-S;
RX MEDLINE=96068664; PubMed=7479849;
RA Fisk G.J., Thummel C.S.;
RT "Isolation, regulation, and DNA-binding properties of three
   Drosophila nuclear hormone receptor superfamily members.",
   Proc. Natl. Acad. Sci. U.S.A. 92:10604-10608 (1995).
RL
CC -1- FUNCTION: BINDS TO NGF1-B RESPONSE ELEMENTS. PLAYS AN IMPORTANT
   ROLE IN LATE STAGES OF EPIDERMAL METAMORPHOSIS.
CC -1- SUBUNIT: FORMS A HETERODIMER WITH USP.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- ALTERNATIVE PRODUCTS:
   Event=Alternative splicing; Named isoforms=2;
   Name=Long;
   IsoId=P49869-1; Sequence=Displayed;
   Name=Short;
   IsoId=P49869-2; Sequence=VSP_003714;
CC -1- TISSUE SPECIFICITY: UBICITOUSLY EXPRESSED IN PRELATOBERN
   EMBRYOS, SPECIALLY IN CENTRAL NERVOUS SYSTEM AND INTSTINAL
   TRACT. HIGHLY EXPRESSED IN THIRD INSTAR LARVAL IMAGINAL DISKS AND
   BRAIN COMPLEXES, BUT NOT IN OVARIES.
CC -1- DEVELOPMENTAL STAGE: LOW LEVELS IN 0-8 HOUR EMBRYOS AND ADULTS.
   HIGHER IN LATE EMBRYOGENESIS AND DURING LARVAL AND PUPAL STAGES.
CC SHORT ISOFORM IS ENRICHED IN PUPAE AND ADULTS, LONG ISOFORM IN
   LARVAE.
CC -1- SIMILARITY: Belongs to the nuclear hormone receptor family. NR4
   subfamily.
CC -----
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CC -----
DR EMBL; X89246; CAA61534.1; -.
DR EMBL; Y15606; CAA75690.1; -.
DR EMBL; AJ002073; CAA05172.1; -.
DR EMBL; AB003667; AAF53914.1; -.
DR EMBL; U36762; AAC46926.1; -.
DR HSSB; P19793; ZNLU.
DR TRANSFAC; T02760; -.
DR FlyBase; FBgn0014859; Hr38.
DR InterPro; IPR000536; Hormone_rec_lig.
DR InterPro; IPR001723; Strdhm_receptor.
DR InterPro; IPR001628; Zn_f_CsteroId.
DR Pfam; PF00104; hormone_rec; 1.
DR Pfam; PF00105; zf-C4; 1.
DR PRINTS; PR00398; STRODHOMNER.
DR PRINTS; PR00047; STROIDINGER.
DR ProDom; PD000035; Zn_f_CsteroId; 1.
DR SMART; SMO0430; HOLY_1.
DR SMART; SMO0399; Zn_f_C4; 1.
DR PROSITE; PS00031; NOCLEAR_RECEPTOR; 1.
KW Receptor; Transcription regulation; DNA-binding; Nuclear protein;
   zinc-finger; Alternative splicing; Developmental protein.
FT znc-finger 744
FT DNA_BIND 744 809
FT          NUCLEAR RECEPTOR-TYPE.
FT ZN_FING 744 764
FT          C4-TYPE.
FT ZN_FING 780 804
FT          C4-TYPE.
FT DOMAIN 188 192
FT          POLY-ALA.
FT DOMAIN 206 218
FT          POLY-ALA.
FT DOMAIN 221 228
FT          POLY-ALA.
FT DOMAIN 258 272
FT          POLY-THR.
FT DOMAIN 294 312
FT          POLY-GLN.
FT DOMAIN 441 462
FT          POLY-GLN.
FT DOMAIN 505 508
FT          POLY-SER.
FT DOMAIN 619 626
FT          POLY-GLN.
FT DOMAIN 661 665
FT          POLY-ALA.
FT VASNPLOC 1 522
           Missing (in isoform Short).
FT /FTid=VSP_003714.

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FT CONFLICT 667 667 V -> VSSPV (IN REF. 4).
FT CONFLICT 685 685 S -> L (IN REF. 1 AND 3).
FT CONFLICT 689 692 STAO -> LGER (IN REF. 2).
FT CONFLICT 697 697 A -> D (IN REF. 2).
FT CONFLICT 702 702 N -> S (IN REF. 2).
FT CONFLICT 1041 1041 S -> R (IN REF. 2).
FT CONFLICT 1064 1064 E -> D (IN REF. 2).
SQ SEQUENCE 1073 AA; 116991 MW; 126A30DAFAC096A CRC64;

Query Match 62.4%; Score 68; DB 1; Length 1073;
Best Local Similarity 73.9%; Pred. No. 2.1;
Matches 17; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

Oy 3 AAAAAAAAAAAAAAAAAAAAAA 25
Db 206 AATAAATPAAAAEAGAAASAAAA 228

RESULT 11
PMXB_HUMAN STANDARD; PRT; 314 AA.
ID PMXB_HUMAN
AC Q99453;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DE 28-FEB-2003 (Rel. 41, Last annotation update)
DE Paired mesoderm homeobox protein 2B (Paired-like homeobox 2B)
DE PHOX2B OR PMX2B.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=97191543; PubMed=9039501;
RA Yokoyama M., Nishi Y., Yoshii J., Okubo K., Matsubara K.;
RT Identification and cloning of neuroblastoma-specific and nerve
RT tissue-specific genes through compiled expression profiles.";
RL DNA Res. 3:311-320(1996).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=20487360; PubMed=11034547;
RA Adachi M., Browne D., Lewis E.J.;
RT "Paired-like homeodomain proteins Phox2a/Arx and Phox2b/NBPHox have
RT similar genetic organization and independently regulate dopamine
RT beta-hydroxylase gene transcription.";
RL DNA Cell Biol. 19:539-554(2000).
RN [3]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=99326521; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [4]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [5]
RP SEQUENCE FROM N.A.
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RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [6]
RP SEQUENCE FROM N.A.
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RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [7]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [8]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
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RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [9]
RP SEQUENCE FROM N.A.
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RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [10]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [11]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [12]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
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RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [13]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
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RX MEDLINE=10395798; PubMed=10395798;
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RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [15]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
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RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [16]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
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RT "Genomic structure and functional characterization of NBPHox (PMX2B),
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RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [17]
RP SEQUENCE FROM N.A.
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RX MEDLINE=10395798; PubMed=10395798;
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RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [18]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
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RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [19]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
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RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [20]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [21]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [22]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [23]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [24]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [25]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [26]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [27]
RP SEQUENCE FROM N.A.
RC TISSUE=Neuroblastoma;
RX MEDLINE=10395798; PubMed=10395798;
RA Yokoyama M., Matcabe H., Nakamura M.;
RT "Genomic structure and functional characterization of NBPHox (PMX2B),
RT a homeodomain protein specific to catecholaminergic cells that is
RT involved in second messenger-mediated transcriptional activation.";
RL Genomics 59:40-50(1999).
RN [28]
RP SEQUENCE FROM
```

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CC -----

DR EMBL; D82344; BA01555.1; -

DR EMBL; AF117979; AAD26698.1; -

DR EMBL; AB015671; BAA82670.1; -

DR PIR; JCS273; JCS273.

DR HSSP; P06601; IFTL.

DR TRANSFAC; T03961; -

DR Genew; HGNC:9143; PHOX2B.

DR MIM; 603851; -

DR GO; GO:0003712; F:transcription cofactor activity; TAS.

DR GO; GO:0003700; F:transcription factor activity; TAS.

DR GO; GO:0007399; P:neurogenesis; TAS.

DR InterPro; IPR001356; Homeobox.

DR InterPro; IPR007104; Paired homeo.

DR Pfam; PF00046; homeobox; 1.

DR Prodom; PD000010; Homeobox; 1.

DR SMART; SM00389; HOX; 1.

DR PROSITE; PS00027; HOMEBOX_1; 1.

DR PROSITE; PS50071; HOMEBOX_2; 1.

DR Homeobox; DNA-binding; Developmental protein; Nuclear protein;

KW Transcription regulation; Activator.

FT DNA BIND 98 157 HOMEBOX.

FT DOMAIN 159 167 POLY-ALA.

FT DOMAIN 212 217 POLY-GLY.

FT DOMAIN 241 260 POLY-ALA.

SO SEQUENCE 314 AA; 31607 MW; 76737F71948B5D81 CRC64;

Query Match 61.5%; Score 67; DB 1; Length 314;

Best Local Similarity 78.3%; Pred. No. 1;

Matches 18; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

OY 3 AAAAAAAAAAAAAAAAAAAAAA 25

DB 244 AAAAAAAAAAAAAAAAAAGLAAA 266

RESULT 12

PMXB_MOUSE STANDARD; PRT; 314 AA.

AC 035690;

DT 15-JUL-1999 (Rel. 38, Last sequence update)

DT 15-JUL-1999 (Rel. 38, Last sequence update)

DT 28-FEB-2003 (Rel. 41, Last annotation update)

DE Paired mesoderm homeobox protein 2B (Paired-like homeobox 2B)

DE (PHOX2B homeodomain protein) (Neuroblastoma Phox) (NBPphox).

OS PHOX2B OR PMX2B.

OS Mus musculus (Mouse).

OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

OX NCBI_TaxID=10090;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=98040559; PubMed=9374403;

RA Paltyn A., Morin X., Cremer H., Goridis C., Brunet J.-F.;

RT "Expression and interactions of the two closely related homeobox genes Phox2a and Phox2b during neurogenesis.";

RL Development 124:4065-4075(1997).

RN [2]

RP SEQUENCE FROM N.A.

RX MEDLINE=99326521; PubMed=10395798;

RA Yokoyama M., Watanabe H., Nakamura M.;

RT "Genomic structure and functional characterization of NBPhox (PMX2B), a homeodomain protein specific to catecholaminergic cells that is involved in second messenger-mediated transcriptional activation.";

RL Genomics 59:40-50(1999).

CC -1- SUBCELLULAR LOCATION: Nuclear (By similarity).

CC -1- SIMILARITY: BELONGS TO THE PAIRED HOMEBOX FAMILY.

CC -----

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CC -----

DR EMBL; Y14493; CAAT483.1; -

DR EMBL; AB015672; BAA82671.1; -

DR HSSP; P06601; IFTL.

DR TRANSFAC; T03976; -

DR MGD; MGI:1100882; Phox2b.

DR InterPro; IPR001356; Homeobox.

DR InterPro; IPR007104; Paired homeo.

DR Pfam; PF00046; homeobox; 1.

DR Prodom; PD000010; Homeobox; 1.

DR SMART; SM00389; HOX; 1.

DR PROSITE; PS00027; HOMEBOX_1; 1.

DR PROSITE; PS50071; HOMEBOX_2; 1.

DR Homeobox; DNA-binding; Developmental protein; Nuclear protein.

FT DNA BIND 98 157 HOMEBOX.

FT DOMAIN 159 167 POLY-ALA.

FT DOMAIN 212 217 POLY-GLY.

FT DOMAIN 241 260 POLY-ALA.

SO SEQUENCE 314 AA; 31621 MW; 40737F71948B595A CRC64;

Query Match 61.5%; Score 67; DB 1; Length 314;

Best Local Similarity 78.3%; Pred. No. 1;

Matches 18; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

OY 3 AAAAAAAAAAAAAAAAAAAAAA 25

DB 244 AAAAAAAAAAAAAAAAAAGLAAA 266

RESULT 13

MAZ_HUMAN STANDARD; PRT; 477 AA.

AC P56270; Q15703; Q99443;

DT 15-JUL-1998 (Rel. 36, Created)

DT 15-JUL-1998 (Rel. 36, Last sequence update)

DT 15-SEP-2003 (Rel. 42, Last annotation update)

DE Myc-associated zinc finger protein (MAZ) (Purine-binding transcription factor) (Pur-1) (ZP87) (ZIF87).

GN MAZ.

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Carnivora; Homiidae; Homo.

OX NCBI_TaxID=9606;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=92366479; PubMed=1502157;

RA Bossone S.A., Asselin C., Patel A.J., Marcu K.B.;

RT "MAZ, a zinc finger protein, binds to c-MYC and C2 gene sequences regulating transcriptional initiation and termination.";

RL Proc. Natl. Acad. Sci. U.S.A. 89:7452-7456(1992).

RN [2]

RP SEQUENCE FROM N.A.

RX MEDLINE=92232709; PubMed=1567856;

RA Pyrc J.J., Moberg K.H., Hall D.J.;

RT "Isolation of a novel cDNA encoding a zinc-finger protein that binds to two sites within the c-myc promoter.";

RL Biochemistry 31:4102-4110(1992).

RN [3]

RP SEQUENCE FROM N.A.

RX MEDLINE=96428591; PubMed=8831693;

RA Tsuboi H., Sakatsume O., Itakura K., Yokoyama K.K.;

RT "Members of the MAZ family: a novel cDNA clone for MAZ from human pancreatic islet cells.";

RL Biochem. Biophys. Res. Commun. 226:801-809(1996).

RN [4]

RP SEQUENCE FROM N.A.

RX MEDLINE=96224025; PubMed=8626793;

RA Parks C.L., Shenk T.;
 RT "The serotonin 1a receptor gene contains a TATA-less promoter that
 RT responds to MAZ and Sp1.";
 RL J. Biol. Chem. 271:4417-4430(1996).
 (5)
 RN SEQUENCE FROM N.A.
 RC TISSUE=Lymphoblastoma;
 RX MEDLINE=98352105; PubMed=9685418;
 RA Song J., Murakami H., Tautsui H., Tang X., Matsumura M., Itakura K.,
 RA Kanazawa I., Sun K., Yokoyama K.K.;
 RT "Genomic organization and expression of a human gene for Myc-
 RT associated zinc finger protein (MAZ).";
 RL J. Biol. Chem. 273:20603-20614(1998).
 CC -1- FUNCTION: MAY FUNCTION AS A TRANSCRIPTION FACTOR WITH DUAL ROLES
 CC IN TRANSCRIPTION INITIATION AND TERMINATION. BINDS TO TWO SITES,
 CC MEAL1 AND MEAL2, WITHIN THE C-MYC PROMOTER HAVING GREATER
 CC AFFINITY FOR THE FORMER. ALSO BINDS TO MULTIPLE G/C-RICH SITES
 CC WITHIN THE PROMOTER OF THE SP1 FAMILY OF TRANSCRIPTION FACTORS.
 CC -1- TISSUE SPECIFICITY: HEART, BRAIN, PLACENTA, LUNG, LIVER, SKELETAL
 CC MUSCLE, AND PANCREAS. SEEMS NOT TO BE EXPRESSED IN KIDNEY.
 CC -1- SIMILARITY: Contains 6 C2H2-type zinc fingers.
 CC -----
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 CC -----
 DR EMBL; M94046; -; NOT ANNOTATED CDS.
 DR EMBL; M93339; -; NOT ANNOTATED CDS.
 DR EMBL; D85311; BA012728.1; ALT_INIT.
 DR EMBL; U33819; AAB04121.1; ALT_INIT.
 DR EMBL; AB017335; BAA33064.1; -;
 DR PIR; A42170; A42170.
 DR TRANSFAC; T00490; -;
 DR TRANSFAC; T02305; -;
 DR GeneW; HGNC:6914; MAZ.
 DR MIM; 600999; -;
 DR GO; GO:0006367; P:transcription initiation from Pol II promoter; TAS.
 DR GO; GO:0006369; P:transcription termination from Pol II promoter; TAS.
 DR InterPro; IPR007087; Znf C2H2.
 DR Pfam; PF00096; zf-C2H2; 5.
 DR ProDom; PD000003; Znf C2H2; 1.
 DR SMART; SM00355; Znf C2H2; 6.
 DR PROSITE; PS00028; ZINC_FINGER_C2H2_1; 5.
 DR PROSITE; PS50157; ZINC_FINGER_C2H2_2; 5.
 KW Transcription regulation; Zinc-finger; Metal-binding; DNA-binding;
 KW RNA-binding; Repeat; Nuclear protein.
 FT ZN_FING 150 212 C2H2-TYPE 1.
 FT ZN_FING 279 301 C2H2-TYPE 2.
 FT ZN_FING 307 329 C2H2-TYPE 3.
 FT ZN_FING 337 360 C2H2-TYPE 4.
 FT ZN_FING 366 388 C2H2-TYPE 5.
 FT ZN_FING 392 413 C2H2-TYPE 6.
 FT DOMAIN 96 108 POLY-ALA.
 FT DOMAIN 133 139 POLY-PRO.
 FT DOMAIN 157 161 POLY-ALA.
 FT DOMAIN 245 249 POLY-GLY.
 FT DOMAIN 435 449 POLY-ALA.
 FT CONFLICT 259 259 MISSING (IN REF. 3).
 FT CONFLICT 401 401 L->M (IN REF. 2 AND 4).
 FT CONFLICT 443 447 MISSING (IN REF. 3).
 SQ SEQUENCE 477 AA; 48607 MW; C04C80F32C3C6825 CRC64;

Query Match 61.5%; Score 67; DB 1; Length 477;
 Best Local Similarity 77.3%; Pred. No. 1.4;

Matches 17; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

4 AAAAAAAAAAAAAAAAAAAAAA 25
 ||| :||| ||||| ||

DB 90 AAAAAAAAAAAAAAAAAAAAAA 111
 RESULT 14
 ID HMD DROAN STANDARD; PRT; 606 AA.
 AC P22544;
 DT 01-AUG-1991 (Rel. 19, Created)
 DT 01-AUG-1991 (Rel. 19, Last sequence update)
 DT 15-SEP-2003 (Rel. 42, Last annotation update)
 DE Homeobox protein B-H1.
 GN B-H1 OR OM(1D).
 OS Drosophila ananassae (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OC NCBI_TaxID=7217;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=91122048; PubMed=1671353;
 RA Tanda S., Corces V.G.;
 RT "Retrotransposon-induced overexpression of a homeobox gene causes
 RT defects in eye morphogenesis in Drosophila.";
 RL EMBL J. 10:407-417(1991).
 CC -1- FUNCTION: Probably involved in eye morphogenesis.
 CC -1- SUBCELLULAR LOCATION: Nuclear (Potential).
 CC -1- SIMILARITY: BELONGS TO THE ANTP HOMEOBOX FAMILY.
 CC -----
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 CC -----
 DR EMBL; X56682; CAA40011.1; -;
 DR PIR; S13367; S13367.
 DR HSSP; P14653; 1872.
 DR TRANSFAC; T03732; -;
 DR FlyBase; FBgn012114; Dana/B-H1.
 DR InterPro; IPR001356; Homeobox.
 DR Pfam; PF00046; homeobox; 1.
 DR PRINTS; PR00024; HOMEOBOX.
 DR ProDom; PD000010; Homeobox; 1.
 DR SMART; SM00389; HOX; 1.
 DR PROSITE; PS00027; HOMEOBOX_1; 1.
 DR PROSITE; PS50071; HOMEOBOX_2; 1.
 KW DNA-binding; Homeobox; Developmental protein; Nuclear protein; Vision.
 FT DOMAIN 23 57 HIS/GLN-RICH (OPA-REPEAT).
 FT DOMAIN 106 124 HIS/GLN-RICH (OPA-REPEAT).
 FT DOMAIN 173 193 HIS/PRO-RICH.
 FT DOMAIN 331 390 HOMEBOX.
 FT DNA_BIND 220 248 ALA-RICH.
 FT DOMAIN 422 434 ALA-RICH.
 FT DOMAIN 450 455 ALA-RICH.
 FT DOMAIN 503 510 ALA-RICH.
 FT DOMAIN 515 521 PRO-RICH.
 SQ SEQUENCE 606 AA; 61735 MW; AA7BB86367370FBB CRC64;

Query Match 61.5%; Score 67; DB 1; Length 606;
 Best Local Similarity 78.3%; Pred. No. 1.7;

Matches 18; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

3 AAAAAAAAAAAAAAAAAAAAAA 25
 ||| :||| ||||| ||

RESULT 15
 HXAD MOUSE STANDARD; PRT; 386 AA.
 ID HXAD MOUSE
 AC Q62424;

DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Homeobox protein Hox-A13 (Hox-1.10).
GN HoxA13 OR Hox-1.10.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxId=10090;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=96259555; PubMed=8673126;
RA Mortlock D.P., Post L.C., Innis J.W.;
RT "The molecular basis of hypodactyly (Hd): a deletion in Hoxa 13 leads
RT to arrest of digital arch formation.";
RL Nat. Genet. 13:284-289(1996).
CC -! FUNCTION: SEQUENCE-SPECIFIC TRANSCRIPTION FACTOR WHICH IS PART OF
CC A DEVELOPMENTAL REGULATORY SYSTEM THAT PROVIDES CELLS WITH
CC SPECIFIC POSITIONAL IDENTITIES ON THE ANTERIOR-POSTERIOR AXIS (BY
CC SIMILARITY).
CC -! SUBCELLULAR LOCATION: Nuclear.
CC -! DISEASE: DEFECTS IN HoxA13 ARE THE CAUSE OF HYPODACTYLY (HD), A
CC CONDITION CHARACTERIZED BY PROFOUND DEFICIENCY OF DIGITAL ARCH
CC STRUCTURES.
CC -! SIMILARITY: BELONGS TO THE ABD-B HOMEBOX FAMILY.
CC -----
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CC or send an email to license@sib-sib.ch).
CC -----
CC EMBL: U59322; AAB03322.1; -.
DR HSSP: P14653; 1872.
DR TRANSPAC: T03337; -.
DR MGD: MGI:96173; Hoxa13.
DR InterPro: IPR001356; Homeobox.
DR Pfam: PF00046; homeobox; 1.
DR ProDom: PD000010; Homeobox; 1.
DR SMART: SMO0389; HOX; 1.
DR PROSITE: PS00027; HOMEBOX_1; 1.
DR PROSITE: PS50071; HOMEBOX_2; 1.
DR Homeobox; DNA-binding; Developmental protein; Nuclear protein;
KW Transcription regulation.
KW DOMAIN 38 51 POLY-ALA.
FT DNA BIND 320 379 HOMEBOX.
FT DOMAIN 52 57 POLY-GLY.
FT DOMAIN 62 66 POLY-ALA.
FT DOMAIN 73 84 POLY-ALA.
FT DOMAIN 101 104 POLY-ALA.
FT DOMAIN 116 133 POLY-ALA.
FT DOMAIN 198 205 POLY-ALA.
SQ SEQUENCE 386 AA; 39566 MW; 2B01DCC9B1951324 CRC64;
Query Match 60.6%; Score 66; DB 1; Length 386;
Best Local Similarity 70.8%; Pred. No. 1.5;
Matches 17; Conservative 2; Mismatches 5; Indels 0; Gaps 0;
QY 2 EAAAAAAAAAAAAAAAAAAAAA 25
Db 111 EAPPSAAAAAAAAAAAAAAAAAS 134

Search completed: January 30, 2004, 00:20:45
Job time : 6.6338 secs

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OM protein - protein search, using sw model

Run on: January 29, 2004, 23:58:52 : Search time 25.8216 Seconds
(without alignments)
249.842 Million cell updates/sec

Title: US-09-461-684C-2
Perfect score: 109
Sequence: 1 CEAATAAATAAATAAATAAATAA 25

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues
Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

SPTREMBL_23:*

- 1: sp_archaea:*
- 2: sp_bacteria:*
- 3: sp_fungi:*
- 4: sp_human:*
- 5: sp_invertebrate:*
- 6: sp_mammal:*
- 7: sp_mhc:*
- 8: sp_organelle:*
- 9: sp_phage:*
- 10: sp_plant:*
- 11: sp_rodent:*
- 12: sp_virus:*
- 13: sp_vertebrate:*
- 14: sp_unclassified:*
- 15: sp_virus:*
- 16: sp_bacteriap:*
- 17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	84	77.1	1038	5 Q8MOW9	Q8MOW9 drosophila
2	84	77.1	2347	5 Q8INH9	Q8INH9 drosophila
3	84	77.1	2451	5 Q9VG05	Q9VG05 drosophila
4	82	75.2	378	11 Q8R089	Q8R089 mus musculus
5	80	73.4	301	11 Q8BJK2	Q8BJK2 mus musculus
6	80	73.4	1354	11 Q9EPW8	Q9EPW8 mus musculus
7	79	72.5	110	11 Q91WV0	Q91WV0 mus musculus
8	79	70.6	332	4 Q81ZU1	Q81ZU1 homo sapien
9	77	70.5	265	10 Q39598	Q39598 chlamydomon
10	76	69.7	665	11 Q90XG2	Q90XG2 mus musculus
11	76	69.7	1452	4 Q9H4A0	Q9H4A0 homo sapien
12	76	69.7	1512	4 Q9H4A1	Q9H4A1 homo sapien
13	75	68.8	221	5 Q9VXD3	Q9VXD3 drosophila
14	72	66.1	997	5 Q9W2J2	Q9W2J2 drosophila
15	71	65.1	206	10 Q39597	Q39597 chlamydomon
16	71	65.1	1216	10 Q9SWH3	Q9SWH3 chlamydomon

17	71	65.1	1787	10 Q9M4X9	Q9M4X9 chlamydomon
18	70	64.2	246	16 Q8G3U2	Q8G3U2 bifidobacte
19	70	64.2	323	4 Q9H782	Q9H782 homo sapien
20	70	64.2	349	4 Q43856	Q43856 homo sapien
21	70	64.2	423	4 Q75400	Q75400 homo sapien
22	70	64.2	452	4 Q75400	Q75400 homo sapien
23	70	64.2	484	11 Q923T4	Q923T4 mus musculus
24	70	64.2	512	4 Q8NAP8	Q8NAP8 homo sapien
25	70	64.2	618	11 Q8BX04	Q8BX04 mus musculus
26	70	64.2	618	11 Q8BX04	Q8BX04 mus musculus
27	70	64.2	1171	3 Q9P3E2	Q9P3E2 neurospora
28	69	63.3	218	11 Q9CS12	Q9CS12 mus musculus
29	69	63.3	512	11 Q8CB12	Q8CB12 mus musculus
30	69	63.3	1222	5 Q8T8L9	Q8T8L9 drosophila
31	69	63.3	1354	5 Q8MPV4	Q8MPV4 drosophila
32	69	63.3	1668	5 Q76930	Q76930 drosophila
33	69	63.3	1669	5 Q9V727	Q9V727 drosophila
34	68	62.4	403	16 Q9JUT2	Q9JUT2 neisseria m
35	68	62.4	543	5 Q9W4F9	Q9W4F9 drosophila
36	68	62.4	545	12 Q9W4F9	Q9W4F9 drosophila
37	68	62.4	1340	16 Q9L1H8	Q9L1H8 tupatia herp
38	68	62.4	2023	5 Q9V5Z9	Q9V5Z9 drosophila
39	68	62.4	2023	5 Q9V5Z9	Q9V5Z9 drosophila
40	67.5	61.9	324	5 Q9N6K2	Q9N6K2 drosophila
41	67.5	61.9	324	5 Q9NGI7	Q9NGI7 drosophila
42	67.5	61.9	324	5 Q9NGI8	Q9NGI8 drosophila
43	67.5	61.9	324	5 Q9N6K3	Q9N6K3 drosophila
44	67.5	61.9	324	5 Q9NGI9	Q9NGI9 drosophila
45	67	61.5	324	5 Q9NGB4	Q9NGB4 drosophila

ALIGNMENTS

RESULT 1

ID	Q8MOW9	PRELIMINARY;	PRT;	1038 AA.
AC	Q8MOW9:			
DT	01-OCT-2002 (TREMBLrel. 22, Created)			
DT	01-OCT-2002 (TREMBLrel. 22, Last sequence update)			
DT	01-MAR-2003 (TREMBLrel. 23, Last annotation update)			
DE	SD05989P (Fragment).			
GN	CG7518.			
OS	Drosophila melanogaster (fruit fly).			
OC	Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;			
OC	Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;			
OC	Ephydroidea; Drosophilidae; Drosophila.			
OX	NCBI_TaxID=7227;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RA	Stapleton W., Brokstein P., Hong L., Agbayani A., Carlson J.,			
RA	Champe M., Chavez C., Dorsett V., Dresnek D., Fafan D., Frise E.,			
RA	George R., Gonzalez M., Guarin H., Krommiller B., Li P., Liao G.,			
RA	Miranda A., Mungall C.J., Nunoo J., Pacleb J., Paragas V., Park S.,			
RA	Patel S., Phouanavong S., Wan K., Yu C., Lewis S.E., Rubin G.M.,			
RA	Celniker S.,			
RL	Submitted (JUN-2002) to the EMBL/GenBank/DBJ databases.			
RL	EMBL; AY122252; AAM52764.1; -			
DR	FlyBase; FBgn0038108; CG7518.			
DR	InterPro; IPR00104; Antifreeze 1.			
DR	InterPro; IPR002965; P rich extensn.			
DR	PRINTS; PR00308; ANTIFREEZE1.			
DR	PRINTS; PR01217; PRICHTEXTENS.			
FT	NON TER			
FT	1			
SC	SEQUENCE 1038 AA; 109059 MW; 80C935A2C6D8A276 CRC64;			

Query Match 77.1%; Score 84; DB 5; Length 1038;
Best Local Similarity 91.3%; Pred. No. 1.1;
Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 3 AAAAAAAAAAAAAAAAAAAAAA 25
Db 112 AAAAAAAAAAAAAAAAAAAAAA 134

RESULT 2
 Q8INH9 PRELIMINARY; PRT; 2347 AA.
 ID Q8INH9;
 AC Q8INH9;
 DT 01-MAR-2003 (TREMBLrel. 23, Created)
 DT 01-MAR-2003 (TREMBLrel. 23, Last sequence update)
 DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
 DE CG7518-PB.
 GN CG7518.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydriidae; Drosophilidae; Drosophila.
 OC NCBI_taxid=7227;
 RP SEQUENCE FROM N.A.
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galie R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.H., Blazer R.G., Champe M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Gabor G.L.,
 RA Abril J.F., Agbayani A., An H.J., Andrews-Pfannkoch C., Baldwin D.,
 RA Ballew R.W., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Bernan B.P., Bhandari D., Bolshakov S.,
 RA Borokova D., Botchan M.R., Bouck J., Brokstein P., Brotlier P.,
 RA Burris K.C., Busan D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablo B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durkin K.J., Evangelista C.C., Ferraz C., Ferrieria S., Fleischmann W.,
 RA Foster C., Gabrielian A.E., Gary N.S., Gelbart W.M., Glaeser K.,
 RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Jaisli D., Houston K.A., Howland T.J., Wei M.H., Ibegwan C.,
 RA Jaisli B., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Lascko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nusser D.R., Pacleb J.M.,
 RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puti V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Svitek R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.Y., Wasserman D.A., Weinstein G.M., Weissenbach J.,
 RA Williams S.M., Woodgett, Worley K.C., Wu D., Yang S., Yao Q.A., Ye J.,
 RA Yeh R.F., Zaveri J.S., Zhan M., Zhang G., Zhu Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhou W., Zhou X., Zhu X., Zhu H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of Drosophila melanogaster.";
 RL Science 287:2185-2195(2000).
 RN (12)
 RP SEQUENCE FROM N.A.
 RA Celniker S.E., Adams M.D., Krommiller B., Wan K.H., Holt R.A.,
 RA Evans C.A., Gocayne J.D., Amanatides P.G., Brandon R.C., Rogers Y.,
 RA Barton J., An H., Baldwin D., Barton J., Beeson K.Y., Busan D.A.,
 RA Carlson J.M., Center A., Champe M., Davenport L.B., Dietz S.M.,
 RA Dodson K., Dorett V., Doup L.E., Doyle C., Drensek D., Fartan D.,
 RA Ferreira S., Frise E., Galie R.F., Gary N.S., George R.A.,
 RA Gonzalez M., Houck J., Hoskins R.A., Hostin D., Howland T.J.,
 RA Ibegwan C., Jaisli M., Kruse D., Li P., Mattei B., Moshrefi A.,
 RA McIntosh T.C., Moy M., Murphy B., Nelson C., Nelson K.A., Nunoo J.,
 RA Pacleb J., Paragay V., Park S., Patel S., Pfeiffer B.,
 RA Phouanavong S., Pittman G.S., Puti V., Richards S., Scheeler F.,
 RA Stapleton M., Strong R., Svitek R., Tector C., Tyler D.,
 RA Williams S.M., Zaveri J.S., Smith H.O., Venter J.C., Rubin G.M.;
 RT "Sequencing of Drosophila melanogaster genome.";

RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RA Misra S., Crosby M.A., Matthews B.B., Bayraktaroglu L., Campbell K.,
 RA Hradecky P., Huang Y., Kaminker J.S., Prochnik S.E., Smith C.D.,
 RA Tupy J.L., Bergman C., Bernan B., Carlson J.W., Celniker S.E.,
 RA Clamp M., Drysdale R., Emmert D., Frise E., de Grey A., Harris N.,
 RA Krommiller B., Marshall B., Millburn G., Richter J., Russo S.,
 RA Searle S.M.J., Smith E., Shu S., Smutnick F., Whitfield E.,
 RA Ashburner M., Gelbart W.M., Rubin G.M., Mungall C.J., Lewis S.E.;
 RT "Annotation of Drosophila melanogaster genome.";
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.
 RA Adams M.D., Celniker S.E., Gibbs R.A., Rubin G.M., Venter J.C.;
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 RN [5]
 RP SEQUENCE FROM N.A.
 RA FlyBase;
 RL Submitted (SEP-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL, AE003598; AAN14338.1; -; 23BF5FC5FCAEA64 CRC64;
 SQ SEQUENCE 2347 AA; 237013 MW; 23BF5FC5FCAEA64 CRC64;
 Query Match 77.1%; Score 84; DB 5; Length 2347;
 Best Local Similarity 91.3%; Pred. No. 2.3;
 Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 3 AAAAAAAAAAAAAAAAAAAAAA 25
 DB 1374 AAAAAAAAAAAAAAAAAAAAAA 1396
 RESULT 3
 Q9VG05 PRELIMINARY; PRT; 2451 AA.
 AC Q9VG05;
 DT 01-MAY-2000 (TREMBLrel. 13, Created)
 DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
 DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
 DE CG7518 protein.
 GN CG7518.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydriidae; Drosophilidae; Drosophila.
 OC NCBI_taxid=7227;
 RP SEQUENCE FROM N.A.
 RC STRAIN=BERKELEY;
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galie R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.H., Blazer R.G., Champe M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Gabor G.L.,
 RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
 RA Ballew R.W., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Bernan B.P., Bhandari D., Bolshakov S.,
 RA Borokova D., Botchan M.R., Bouck J., Brokstein P., Brotlier P.,
 RA Burris K.C., Busan D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablo B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durkin K.J., Evangelista C.C., Ferraz C., Ferrieria S., Fleischmann W.,
 RA Foster C., Gabrielian A.E., Gary N.S., Gelbart W.M., Glaeser K.,
 RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Jaisli D., Houston K.A., Howland T.J., Wei M.H., Ibegwan C.,
 RA Jaisli B., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Lascko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,

RA Merkulyov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nusser D.R., Pacle J.M.,
 RA Palazzo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Svitzkas R., Tecor C., Turner E., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weissbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of *Drosophila melanogaster*.";
 RL Science 287:2185-2195(2000).
 DR EMBL; AF003698; AAF54888.2;..
 DR FlyBase; FBgn0038108; CG7518.
 DR InterPro; IPR001005; MYB_DNA_binding.
 DR PROSITE; PS00037; MYB 1; 1.
 SQ SEQUENCE 2451 AA; 266959 MW; 088A2293F27481E2 CRC64;

Query Match 77.1%; Score 84; DB 5; Length 2451;
 Best Local Similarity 91.3%; Pred. No. 2.4;
 Matches 21; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Oy 3 AAAAAAAAAAAAAAAAAAAAAA 25
 Db 1374 AAAAAAAAAAAAAAAAAAAAAA 1396

RESULT 4

O8R089 PRELIMINARY; PRT; 378 AA.

ID O8R089; PRELIMINARY; PRT; 378 AA.
 AC O8R089; PRELIMINARY; PRT; 378 AA.
 DT 01-JUN-2002 (TRENBLrel. 21, Created)
 DT 01-JUN-2002 (TRENBLrel. 21, Last sequence update)
 DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
 DE Similar to hypothetical protein FLJ11618.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 NC NCB1_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Colon;
 RA Straubeberg R.;
 RL Submitted (Apr-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL; BC027193; AAH27193.1;..
 KW Hypothetical protein.
 SQ SEQUENCE 378 AA; 39456 MW; 4C3FAF0DAAC29E69 CRC64;

Query Match 75.2%; Score 82; DB 11; Length 378;
 Best Local Similarity 87.5%; Pred. No. 0.73;
 Matches 21; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Oy 2 EAAAAAAAAAAAAAAAAAAAAA 25
 Db 222 EAAAAAAAAAAAAAAAAAAAAA 245

RESULT 5

O8BUK2 PRELIMINARY; PRT; 301 AA.

ID O8BUK2; PRELIMINARY; PRT; 301 AA.
 AC O8BUK2; PRELIMINARY; PRT; 301 AA.
 DT 01-MAR-2003 (TRENBLrel. 23, Created)
 DT 01-MAR-2003 (TRENBLrel. 23, Last sequence update)
 DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
 DE Cell division cycle 2-like 5 (Fragment).
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 NC NCB1_TaxID=10090;
 RN [1]

RP SEQUENCE FROM N.A.
 RC STRAIN=CS7BL/6J; TISSUE=Body;
 RX MEDLINE=22354683; PubMed=12466851;
 RA The PANTOM Consortium,
 RA the RIKEN Genome Exploration Research Group Phase I & II Team;
 RT "Analysis of the mouse transcriptome based on functional annotation of
 RT 60,770 full-length cDNAs.";
 RL Nature 420:563-573(2002).
 DR EMBL; AK083577; BAC38957.1;..
 FT NON TER 1
 SQ SEQUENCE 301 AA; 32269 MW; A6CF891DBE25E09E CRC64;

Query Match 73.4%; Score 80; DB 11; Length 301;
 Best Local Similarity 83.3%; Pred. No. 0.93;
 Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Oy 2 EAAAAAAAAAAAAAAAAAAAAA 25
 Db 74 EAAAAAAAAAAAAAAAAAAAAA 97

RESULT 6

O9EPW8 PRELIMINARY; PRT; 1354 AA.

ID O9EPW8; PRELIMINARY; PRT; 1354 AA.
 AC O9EPW8; PRELIMINARY; PRT; 1354 AA.
 DT 01-MAR-2001 (TRENBLrel. 16, Created)
 DT 01-MAR-2001 (TRENBLrel. 16, Last sequence update)
 DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
 DE Nischcharin.
 GN NISCH.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 NC NCB1_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ICR outbred; TISSUE=Brain;
 RX MEDLINE=20571837; PubMed=11121431;
 RA Alahari S.K., Lee J.W., Juliano R.L.;
 RT "Nischcharin, a Novel Protein That Interacts with the Integrin alphas
 RT Subunit and Inhibits Cell Migration.";
 RL J. Cell Biol. 151:1141-1154(2000).
 DR EMBL; AF153344; AAG42100.1;..
 DR MED; MGI:1928323; Nisch.
 DR InterPro; IPR001128; Cytochrome_P450.
 DR InterPro; IPR001611; LRR.
 DR InterPro; IPR007092; LRR_SDS22.
 DR Pfam; PF00560; LRR; 5.
 DR PRINTS; PRO0019; LEURICRPT.
 DR PROSITE; PS00086; CYTOCHROME_P450; 1.
 DR PROSITE; PS50504; LRR_SDS22; 1.
 SQ SEQUENCE 1354 AA; 148060 MW; 01BD676FDC1A19247 CRC64;

Query Match 73.4%; Score 80; DB 11; Length 1354;
 Best Local Similarity 83.3%; Pred. No. 3.4;
 Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Oy 2 EAAAAAAAAAAAAAAAAAAAAA 25
 Db 837 EAPAAAEAPAAAEAPAAAEAPAA 860

RESULT 7

O91WMO PRELIMINARY; PRT; 110 AA.

ID O91WMO; PRELIMINARY; PRT; 110 AA.
 AC O91WMO; PRELIMINARY; PRT; 110 AA.
 DT 01-DEC-2001 (TRENBLrel. 19, Created)
 DT 01-DEC-2001 (TRENBLrel. 19, Last sequence update)
 DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
 DE Hypothetical 10.7 kDa protein.
 GN A1591529.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

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OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Salivary gland;
RA Strausberg R.;
RL Submitted (Aug-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC012681; AAH12681.1; -
DR MGD; MGI:2139008; A1591529.
DR InterPro; IPR001014; Antifreeze_1.
DR InterPro; IPR01859; Ribosomal_P2.
DR PRINTS; PR00308; ANTIFREEZE1.
DR PRINTS; PR00456; RIBOSOMALP2.
KW Hypothetical protein.
SQ SEQUENCE 110 AA; 10662 MW; 0581D2635F87EAB CRC64;

Query Match
Best Local Similarity 72.5%; Score 79; DB 11; Length 110;
Matches 20; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 2 EAAAAAAAAAAAAAAAAAAAAA 25
DB 23 DAAAAAAAAAAAAAAAAAAAAA 46

RESULT 8
Q81ZU1 PRELIMINARY; PRT; 332 AA.
ID Q81ZU1;
AC Q81ZU1;
DT 01-MAR-2003 (TREMBLrel. 23, Created)
DT 01-MAR-2003 (TREMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Family with sequence similarity 9, member A.
GN FAM9A.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=2202142; PubMed=12213195;
RA Martinez-Garay I., Jablonka S., Sutarajova M., Steurnagel P., Gal A.,
RA Kutsche K.;
RT "A New Gene Family (FAM9) of Low-Copy Repeats in Xp22.3 Expressed
RT Exclusively in Testis: Implications for Recombinations in This
RT Region.";
RL Genomics 80:259-267(2002).
DR EMBL; AF494343; AAN07162.1; -
SQ SEQUENCE 332 AA; 37339 MW; 92F22EG36038229C CRC64;

Query Match
Best Local Similarity 72.5%; Score 79; DB 4; Length 332;
Matches 20; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 EAAAAAAAAAAAAAAAAAAAAA 25
DB 190 EAEFEAAAAAAAAAAAAAAAAA 213

RESULT 9
Q39598 PRELIMINARY; PRT; 265 AA.
ID Q39598;
AC Q39598;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
DE Cgcr-4 product (Fragment).
GN CGCR-4.
OS Chlamydomonas reinhardtii.
OC Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
OC Chlamydomonadaceae; Chlamydomonas.
OX NCBI_TaxID=3055;
RN [1]

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RP SEQUENCE FROM N.A.
RX MEDLINE=92119224; PubMed=1731966;
RA Wakarchuk W.W., Muller P.W., Beck C.F.;
RT "Two GC-rich DNA elements of Chlamydomonas reinhardtii with complex
RT arrangements of directly repeated sequence motifs.";
RL Plant Mol. Biol. 18:143-146(1992).
DR EMBL; X17208; CAA35080.1; -
FT NON TER
SQ SEQUENCE 265 AA; 26216 MW; B35318B7377CF782 CRC64;

Query Match
Best Local Similarity 70.6%; Score 77; DB 10; Length 265;
Matches 19; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 3 AAAAAAAAAAAAAAAAAAAAAA 25
DB 154 AAAAAAAAAAAKARVAAEAA 176

RESULT 10
Q9QXG2 PRELIMINARY; PRT; 665 AA.
ID Q9QXG2;
AC Q9QXG2;
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
DE Choroideremia protein.
GN CHM.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BALB/c;
RA van den Hurk J.A., Huber I., van de Pol T.J., Cremers F.P.;
RT "Cloning and sequencing of the mouse choroideremia gene.";
RL Submitted (DEC-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF218084; AAF25478.1; -
DR MGD; MGI:892979; Cmm.
DR InterPro; IPR02005; Rab_GDI_REP.
DR Pfam; PF00996; GDI_1.
DR PRINTS; PR00891; RABGDIREP.
SQ SEQUENCE 665 AA; 73976 MW; FF71A74AD3FBDE0A CRC64;

Query Match
Best Local Similarity 69.7%; Score 76; DB 11; Length 665;
Matches 19; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 2 EAAAAAAAAAAAAAAAAAAAAA 25
DB 132 EAEAEAAEAETEAEEAAEAA 155

RESULT 11
Q9H4A0 PRELIMINARY; PRT; 1452 AA.
ID Q9H4A0;
AC Q9H4A0;
DT 01-MAR-2001 (TREMBLrel. 16, Created)
DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE CDC2L5 protein kinase.
GN CDC2L5.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Placenta;
RA Marques F., Moreau J.L., Peaucellier G., Lozano J.C., Schatt P.,
RA Picard A., Callebaut I., Perre E., Genevriere A.M.;
RT "A new subfamily of high molecular mass CDC2-related kinases with

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RT PITAI/VRE."
 RL Biochem. Biophys. Res. Commun. 279:832-837(2001).
 CC -I- SIMILARITY: BELONGS TO THE SER/THR FAMILY OF PROTEIN KINASES.
 DR EMBL; AJ297710; CAC10401.1; ...
 DR HSP; F24941; 1BDH.
 DR InterPro; IPR000719; Prot_kinase.
 DR InterPro; IPR002290; Ser_thr_kinase.
 DR Pfam; PF00069; pkinase; 1.
 DR ProDom; PD000001; Prot_kinase; 1.
 DR SMART; SM00220; S_TKC; 1.
 DR PROSITE; PS00107; PROTEIN_KINASE_ATP; 1.
 DR PROSITE; PS00011; PROTEIN_KINASE_DOM; 1.
 DR PROSITE; PS00108; PROTEIN_KINASE_ST; 1.
 KW ATP-binding; Kinase; Serine/threonine-protein kinase; Transferase.
 SQ SEQUENCE 1452 AA; 158480 MW; C7ED072368B439CB CRC64;

Query Match 69.7%; Score 76; DB 4; Length 1452;
 Best Local Similarity 79.2%; Pred. No. 8.6;
 Matches 19; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 2 EAAAAAAAAAAAAAAAAAAAAA 25
 DB 463 EAAARAAAKAAEATKAAEAAAKA 486

RESULT 12

Q9H4A1 PRELIMINARY; PRT; 1512 AA.
 ID Q9H4A1;
 AC Q9H4A1;
 DT 01-MAR-2001 (TrEMBLrel. 16, Created)
 DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE CDC2L5 protein kinase.
 GN CDC2L5.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=placenta;
 RA Marques F., Moreau J.L., Peaucellier G., Lozano J.C., Schatt P.,
 RA Picard A., Callebaut I., Perre E., Genevriere A.M.;
 RT "A new subfamily of high molecular mass CDC2-related kinases with
 PITAI/VRE."
 RL Biochem. Biophys. Res. Commun. 279:832-837(2001).
 CC -I- SIMILARITY: BELONGS TO THE SER/THR FAMILY OF PROTEIN KINASES.
 DR EMBL; AJ297709; CAC10400.1; ...
 DR HSP; P24941; 1BDH.
 DR InterPro; IPR000719; Prot_kinase.
 DR InterPro; IPR002290; Ser_thr_kinase.
 DR Pfam; PF00069; pkinase; 1.
 DR ProDom; PD000001; Prot_kinase; 1.
 DR SMART; SM00220; S_TKC; 1.
 DR PROSITE; PS00107; PROTEIN_KINASE_ATP; 1.
 DR PROSITE; PS00011; PROTEIN_KINASE_DOM; 1.
 DR PROSITE; PS00108; PROTEIN_KINASE_ST; 1.
 KW ATP-binding; Kinase; Serine/threonine-protein kinase; Transferase.
 SQ SEQUENCE 1512 AA; 164969 MW; 2838BD553DB57650 CRC64;

Query Match 69.7%; Score 76; DB 4; Length 1512;
 Best Local Similarity 79.2%; Pred. No. 8.9;
 Matches 19; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 2 EAAAAAAAAAAAAAAAAAAAAA 25
 DB 463 EAAARAAAKAAEATKAAEAAAKA 486

RESULT 13
 Q9VXD3 PRELIMINARY; PRT; 221 AA.
 AC Q9VXD3;

DT 01-MAY-2000 (TrEMBLrel. 13, Created)
 DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
 DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
 DE CG13012 protein.
 GN CG13012.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=BERKELEY;
 RX MEDLINE=20196006; PubMed=10731132;

RA Adams M.D., Gelinker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galie R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutton G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.-H.C., Blazej R.G., Champe M., Pfeiffer B.D.,
 RA Man K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
 RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
 RA Bailew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Bernan B.P., Bhandari D., Bolshakov S.,
 RA Borkova D., Botchan M.R., Bouck J., Brokstein P., Brotlier P.,
 RA Burtis K.C., Busam D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport I.B., Davies P.,
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
 RA Foster C., Gabrieli A.E., Gary N.S., Gelbart W.M., Glaeser K.,
 RA Glodek A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.V., Wei M.-H., Ibegwam C.,
 RA Jaitani B., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Lascko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merkulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nusken D.R., Pacle J.M.,
 RA Palazolo M., Pittman G.S., Pan S., Pollard J., Part V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Svitskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.-Y., Wassarman D.A., Weinstein G.M., Weissbach J.,
 RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
 RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao O., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of Drosophila melanogaster."
 RL Science 287:2185-2195(2000).
 DR EMBL; AE003503; AAF48641.1; ...
 DR FLYbase; FBgn0030769; CG13012.
 SQ SEQUENCE 221 AA; 22987 MW; A1B95919B167C5E2 CRC64;

Query Match 68.8%; Score 75; DB 5; Length 221;
 Best Local Similarity 81.8%; Pred. No. 2.1;
 Matches 18; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 4 AAAAAAAAAAAAAAAAAAAAAA 25
 DB 6 AAAAAAAAAAATVAAATVAAAAA 27

RESULT 14
 Q9W2J2 PRELIMINARY; PRT; 997 AA.
 ID Q9W2J2;
 AC Q9W2J2;
 DT 01-MAY-2000 (TrEMBLrel. 13, Created)
 DT 01-OCT-2002 (TrEMBLrel. 22, Last sequence update)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE CG18375 protein.

Fri Jan 30 06:18:16 2004

us-09-461-684C-2.rpt

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Search completed: January 30, 2004, 00:24:39
Job time : 27.8216 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 30, 2004, 00:13:12 ; Search time 10.2535 Seconds
(without alignments)
225.098 Million cell updates/sec

Title: US-09-461-684C-3

Perfect score: 143

Sequence: 1 GGLFGAIGAFIENGWEGMIDGWYG 24

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283308 seqs, 96168682 residues

Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :
1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	134	93.7	550	1 HMTV52	hemagglutinin prec
2	134	93.7	550	1 HMTV53	hemagglutinin prec
3	134	93.7	550	1 HMTV77	hemagglutinin prec
4	134	93.7	550	1 HMTV80	hemagglutinin prec
5	134	93.7	550	1 HMTV33	hemagglutinin prec
6	134	93.7	550	1 HMTV89	hemagglutinin prec
7	134	93.7	550	1 HMTV21	hemagglutinin prec
8	134	93.7	550	1 HMTV98	hemagglutinin prec
9	134	93.7	550	1 HMTV15	hemagglutinin prec
10	134	93.7	550	2 JOL153	hemagglutinin prec
11	134	93.7	550	2 JOL154	hemagglutinin prec
12	134	93.7	550	2 JOL155	hemagglutinin prec
13	134	93.7	566	1 HMTV8	hemagglutinin prec
14	134	93.7	566	1 HMTV8A	hemagglutinin prec
15	134	93.7	566	1 HMTV8M	hemagglutinin prec
16	134	93.7	566	1 HMTV8U	hemagglutinin prec
17	133	93.0	561	1 HMTV49	hemagglutinin prec
18	133	93.0	561	1 HMTV84	hemagglutinin prec
19	132	92.3	565	1 HMTV81	hemagglutinin prec
20	132	92.3	565	1 HMTV83	hemagglutinin prec
21	132	92.3	566	1 HMTV6	hemagglutinin prec
22	132	92.3	567	1 HMTV6	hemagglutinin prec
23	131	91.6	362	2 S38637	hemagglutinin - in
24	131	91.6	550	1 HMTV86	hemagglutinin prec
25	131	91.6	560	1 HMTV77	hemagglutinin prec
26	131	91.6	565	1 HMTV82	hemagglutinin prec
27	131	91.6	565	1 HMTV84	hemagglutinin prec
28	131	91.6	565	1 HMTV85	hemagglutinin prec
29	131	91.6	565	1 HMTV86	hemagglutinin prec

30	131	91.6	565	1 HMTV87	hemagglutinin prec
31	131	91.6	565	1 HMTV88	hemagglutinin prec
32	131	91.6	565	1 HMTV89	hemagglutinin prec
33	131	91.6	565	1 HMTV8	hemagglutinin prec
34	131	91.6	565	1 HMTV8	hemagglutinin prec
35	131	91.6	565	2 S33703	hemagglutinin - in
36	131	91.6	570	1 A45591	hemagglutinin prec
37	131	91.6	570	2 S22013	hemagglutinin prec
38	131	91.6	570	2 S22014	hemagglutinin prec
39	131	91.6	570	2 S22015	hemagglutinin prec
40	131	91.6	570	2 S22016	hemagglutinin prec
41	131	91.6	570	2 S22017	hemagglutinin prec
42	131	91.6	570	2 S22018	hemagglutinin prec
43	131	91.6	570	2 S22020	hemagglutinin prec
44	131	91.6	570	2 S22021	hemagglutinin prec
45	131	91.6	570	2 S22029	hemagglutinin prec

ALIGNMENTS

RESULT 1
HMTV52 hemagglutinin precursor - Influenza A virus (strain A/swine/126/82) (fragment)
C:Species: influenza A virus
C:Date: 31-Mar-1989 #sequence_revision 31-Mar-1989 #text_change 18-Sep-1998
C:Accession: A29971
R:Kida, H.; Shortridge, K.F.; Webster, R.G.
Virology 162, 160-166, 1988
A:Title: Origin of the hemagglutinin gene of H3N2 influenza viruses from pigs in China.
A:Reference number: A94370; PMID:88101364; PMID:3336940
A:Accession: A29971
A:Molecule type: genomic RNA
A:Residues: 1-550 <KID>
A:Cross-references: GB:M19056; NID:g324208
A>Note: the sequence in Genbank entry FLAHAP, release 106, (PID:g324209) differs from tl
C:Genetics:
A:Map position: segment 4
C:Superfamily: Influenza virus hemagglutinin
C:Keywords: glycoprotein, hemagglutinin, homotrimer, lipoprotein, thiolester bond
F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>
F:130-550/Product: hemagglutinin HA2 #status predicted <HA2>
F:520-556/Product: transmembrane #status predicted <TM1>
F:8-22,38,165,285,483/Binding site: carbohydrate (asn) (covalent) #status predicted
F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted
F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted
Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1,4e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Oy 2 GGLFGAIGAFIENGWEGMIDGWYG 24
DB 330 GGLFGAIGAFIENGWEGMIDGWYG 352
RESULT 2
HMTV53 hemagglutinin precursor - Influenza A virus (strain A/swine/81/78) (fragment)
C:Species: influenza A virus
C:Date: 31-Mar-1989 #sequence_revision 31-Mar-1989 #text_change 18-Sep-1998
C:Accession: B29971
R:Kida, H.; Shortridge, K.F.; Webster, R.G.
Virology 162, 160-166, 1988
A:Title: Origin of the hemagglutinin gene of H3N2 influenza viruses from pigs in China.
A:Reference number: A94370; PMID:88101364; PMID:3336940
A:Accession: B29971
A:Molecule type: genomic RNA
A:Residues: 1-550 <KID>
A:Cross-references: GB:M19057; NID:g324210
A>Note: the sequence in Genbank entry FLAHAPB, release 106, (PID:g324211) differs from tl
C:Genetics:
A:Map position: segment 4

C:Superfamily: influenza virus hemagglutinin
C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond
F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>
F:300-550/Product: hemagglutinin HA2 #status predicted <HA2>
F:520-536/Domain: transmembrane #status predicted <TM1>
F:8,22,38,165,285,483/Binding site: carbohydrate (asn) (covalent) #status predicted
F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted
F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.4e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMIDGWYG 24
DB 330 GLFGAIAGFIENGWEGMIDGWYG 352

RESULT 3
HMI177
hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/5/77) (fragment)

N:Contains: hemagglutinin HA1; hemagglutinin HA2
C:Species: influenza A virus
C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 16-Jul-1999
C:Accession: A27813
R:Kida, H.; Kawoka, Y.; Naeve, C.W.; Webster, R.G.
Virology 159, 109-119, 1987

A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.

A:Reference number: A94363; PMID:87265458; PMID:2440178

A:Accession: A27813

A:Molecule type: genomic RNA

A:Residues: 1-550 <KID>

A:Cross-references: GB:M16737; NID:G324081; PID:AAA3143.1; PID:G324082

C:Genetics:

A:Map position: segment 4

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>

F:520-536/Domain: transmembrane #status predicted <TM1>

F:8,22,38,165,285,483/Binding site: carbohydrate (asn) (covalent) #status predicted

F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted

F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.4e-10;

Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMIDGWYG 24
DB 330 GLFGAIAGFIENGWEGMIDGWYG 352

RESULT 4
HMI180
hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/8/80) (fragment)

N:Contains: hemagglutinin HA1; hemagglutinin HA2
C:Species: influenza A virus
C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 18-Sep-1998
C:Accession: B27813

R:Kida, H.; Kawoka, Y.; Naeve, C.W.; Webster, R.G.
Virology 159, 109-119, 1987

A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.

A:Reference number: A94363; PMID:87265458; PMID:2440178

A:Accession: B27813

A:Molecule type: genomic RNA

A:Residues: 1-550 <KID>

A:Cross-references: GB:M16738; NID:G324083

A:Note: the translation in Fig. 2 is inconsistent with the nucleotide sequence in Fig. 1

C:Genetics:

A:Map position: segment 4

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>

F:520-536/Domain: transmembrane #status predicted <TM1>

F:8,22,38,165,285,483/Binding site: carbohydrate (asn) (covalent) #status predicted

F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted

F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.4e-10;

Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMIDGWYG 24
DB 330 GLFGAIAGFIENGWEGMIDGWYG 352

RESULT 5
HMI173
hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/33/80) (fragment)

N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: influenza A virus

C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 16-Jul-1999

C:Accession: C27813

R:Kida, H.; Kawoka, Y.; Naeve, C.W.; Webster, R.G.

Virology 159, 109-119, 1987

A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.

A:Reference number: A94363; PMID:87265458; PMID:2440178

A:Accession: C27813

A:Molecule type: genomic RNA

A:Residues: 1-550 <KID>

A:Cross-references: GB:M16739; NID:G324085; PID:AAA3145.1; PID:G324086

C:Genetics:

A:Map position: segment 4

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>

F:520-536/Domain: transmembrane #status predicted <TM1>

F:8,22,38,165,285,483/Binding site: carbohydrate (asn) (covalent) #status predicted

F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted

F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.4e-10;

Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMIDGWYG 24
DB 330 GLFGAIAGFIENGWEGMIDGWYG 352

RESULT 6
HMI189

hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/7/82) (fragment)

N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: influenza A virus

C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 16-Jul-1999

C:Accession: D27813

R:Kida, H.; Kawoka, Y.; Naeve, C.W.; Webster, R.G.

Virology 159, 109-119, 1987

A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.

A:Reference number: A94363; PMID:87265458; PMID:2440178

A:Accession: D27813

A:Molecule type: genomic RNA

A:Residues: 1-550 <KID>

A:Cross-references: GB:M16740; NID:G324087; PID:AAA3146.1; PID:G324088

C:Genetics:

A:Map position: segment 4

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>

F:520-536/Domain: transmembrane #status predicted <TM1>

F:8,22,38,165,285,483/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted
F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1,4e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAGFIENGWEGMIDGMYG 24
Db 330 GLFGAIAGFIENGWEGMIDGMYG 352

RESULT 7

HMIIV21
hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/21/62) (fragment)
N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: influenza A virus
C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 18-Sep-1998
C:Accession: E27813
R:Kida, H.; Kawoka, Y.; Naeve, C.W.; Webster, R.G.
Virology 159, 109-119, 1987

A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.
A:Reference number: A94363; MUID:87265458; PMID:2440178

A:Accession: E27813
A:Molecule type: genomic RNA
A:Residues: 1-550 <KID>

A:Cross-references: GB:M16741; NID:9324089
C:Genetics:

A:Map position: segment 4

C:Superfamily: influenza virus hemagglutinin
C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>

F:520-536/Domains: transmembrane #status predicted <TM1>

F:7,8,22,38,165,285,483/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1,4e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAGFIENGWEGMIDGMYG 24
Db 330 GLFGAIAGFIENGWEGMIDGMYG 352

RESULT 8

HMIIV98
hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/9/85) (fragment)

N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: influenza A virus
C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 18-Sep-1998
C:Accession: F27813

R:Kida, H.; Kawoka, Y.; Naeve, C.W.; Webster, R.G.
Virology 159, 109-119, 1987

A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.
A:Reference number: A94363; MUID:87265458; PMID:2440178

A:Accession: F27813

A:Molecule type: genomic RNA
A:Residues: 1-550 <KID>

A:Cross-references: GB:M16742; NID:9324091

C:Genetics:

A:Map position: segment 4

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>

F:520-536/Domains: transmembrane #status predicted <TM1>

F:8,22,38,165,285,483/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted
F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1,4e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAGFIENGWEGMIDGMYG 24
Db 330 GLFGAIAGFIENGWEGMIDGMYG 352

RESULT 9

HMIIV15

hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/10/85) (fragment)
N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: influenza A virus
C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 16-Jul-1999
C:Accession: G27813
R:Kida, H.; Kawoka, Y.; Naeve, C.W.; Webster, R.G.
Virology 159, 109-119, 1987

A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.
A:Reference number: A94363; MUID:87265458; PMID:2440178

A:Accession: G27813

A:Molecule type: genomic RNA
A:Residues: 1-550 <KID>

A:Cross-references: GB:M16743; NID:9324093; PIDN:AAA43149.1; PID:9324094
C:Genetics:

A:Map position: segment 4

C:Superfamily: influenza virus hemagglutinin
C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>

F:520-536/Domains: transmembrane #status predicted <TM1>

F:8,22,38,165,285,483/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted
F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1,4e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAGFIENGWEGMIDGMYG 24
Db 330 GLFGAIAGFIENGWEGMIDGMYG 352

RESULT 10

QJ1153

hemagglutinin precursor - influenza A virus (strain A/duck/Hong Kong/7/75) (fragment)
N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: influenza A virus
C:Date: 31-Mar-1992 #sequence_revision 31-Mar-1992 #text_change 20-Jun-2000
C:Accession: U01153

R:Yaounda, J.; Shortridge, K.F.; Shimizu, Y.; Kida, H.
J. Gen. Virol. 72, 2007-2010, 1991

A:Title: Molecular evidence for a role of domestic ducks in the introduction of avian H3
A:Reference number: QJ1153; MUID:91341491; PMID:1875195

A:Accession: U01153

A:Molecule type: genomic RNA
A:Residues: 1-550 <YAS>

A:Cross-references: GB:D00929; NID:9221279; PIDN:BA00769.1; PID:9221280
A:Note: the authors translated the codon GGG for residue 218 as Glu

A:Note: residues 528-532 are not shown in this publication

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; homotrimer

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-545/Product: hemagglutinin HA2 #status predicted <HA2>

F:8,22,38,165,285,483/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 2; Length 550;
Best Local Similarity 100.0%; Pred. No. 1,4e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAGFIENGWEGMIDGMYG 24
|||||

Db 330 GLFGAIAAGFIENGWEGMIDGMYG 352

RESULT 11

QJ01154

hemagglutinin precursor - influenza A virus (strain A/goose/Hong Kong/10/76) (fragment)
N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Date: 31-Mar-1992 #sequence_revision 31-Mar-1992 #text_change 20-Jun-2000

C:Accession: JQ1154

R:Yaounda, U.; Shortridge, K.F.; Shimizu, Y.; Kida, H.

J. Gen. Virol. 72, 2007-2010, 1991

A:Title: Molecular evidence for a role of domestic ducks in the introduction of avian H3

A:Reference number: JQ1153; MUID:91341491; PMID:1875195

A:Accession: JQ1154

A:Molecule type: genomic RNA

A:Residues: 1-550 <YAS>

A:Cross-references: GB:D00930; NID:g221273; PIDN:BA00770.1; PID:g221274

A:Note: the authors translated the codon GCG for residue 218 as Glu

A:Note: residues 528-532 are not shown in this publication

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; homotrimer

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-545/Product: hemagglutinin HA2 #status predicted <HA2>

F:8,22,38,165,285,483/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 2; Length 550;

Best Local Similarity 100.0%; Pred. No. 1,4e-10;

Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAAGFIENGWEGMIDGMYG 24

Db 330 GLFGAIAAGFIENGWEGMIDGMYG 352

RESULT 12

QJ01155

hemagglutinin precursor - influenza A virus (strain A/duck/Hong Kong/64/76) (fragment)
N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: Influenza A virus

C:Date: 31-Mar-1992 #sequence_revision 31-Mar-1992 #text_change 20-Jun-2000

C:Accession: JQ1155

R:Yaounda, U.; Shortridge, K.F.; Shimizu, Y.; Kida, H.

J. Gen. Virol. 72, 2007-2010, 1991

A:Title: Molecular evidence for a role of domestic ducks in the introduction of avian H3

A:Reference number: JQ1153; MUID:91341491; PMID:1875195

A:Accession: JQ1155

A:Molecule type: genomic RNA

A:Residues: 1-550 <YAS>

A:Cross-references: GB:D00931; NID:g221277; PIDN:BA00771.1; PID:g221278

A:Note: the authors translated the codon GCG for residue 218 as Glu, GCC for residue 538

A:Note: residues 528-532 are not shown in this publication

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; homotrimer

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-545/Product: hemagglutinin HA2 #status predicted <HA2>

F:8,22,38,165,285,483/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 2; Length 550;

Best Local Similarity 100.0%; Pred. No. 1,4e-10;

Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAAGFIENGWEGMIDGMYG 24

Db 330 GLFGAIAAGFIENGWEGMIDGMYG 352

RESULT 13

HM1IVH

hemagglutinin precursor - influenza A virus

C:Species: Influenza A virus

C:Date: 28-Feb-1981 #sequence_revision 28-Feb-1981 #text_change 22-Oct-1999

C:Accession: A93705; A93233; A04051; A93231; A94441

R:Boch, G.W.; Sleight, M.J.

Nucleic Acids Res. 8, 2561-2575, 1980

A:Title: Complete nucleotide sequence of the haemagglutinin gene from a human influenza

A:Reference number: A93705; MUID:81053698; PMID:6253883

A:Accession: A93705

A:Molecule type: genomic RNA

A:Residues: 1-566 <BOT>

A:Cross-references: GB:V01103

A:Experimental source: strain A/NT/60/68/29C

A:Note: human influenza strain A/NT/60/68/29C is a laboratory-isolated variant of A/NT/60

R:Dopheide, T.A.; Ward, C.W.

PNAS Lett. 110, 181-183, 1980

A:Title: The disulfide bonds of a Hong Kong influenza virus hemagglutinin.

A:Reference number: A91276; MUID:80179105; PMID:6768586

A:Contents: annotation; disulfide bonds

R:Geething, M.J.; Bye, J.; Skehel, J.; Waterfield, M.

Nature 287, 301-306, 1980

A:Title: Cloning and DNA sequence of double-stranded copies of haemagglutinin genes from

A:Reference number: A93233; MUID:81030852; PMID:7421990

A:Accession: A93233

A:Molecule type: genomic RNA

A:Residues: 1-24, 'S', 26, 'D', 28-159, 'G', 161-197, 'I', 199-241, 'L', 243-249 <GET>

A:Experimental source: strain X-31(H3)

C:Superfamily: influenza virus hemagglutinin

C:Keywords: hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-16/DNA: signal sequence #status predicted <SIG>

F:17-344/Product: hemagglutinin HA1 #status predicted <HA1>

F:346-566/Product: hemagglutinin HA2 #status predicted <HA2>

F:536-552/DNA: transmembrane #status predicted <TM1>

F:30-482, 68-293, 80-92, 155-489, 293-321/Disulfide bonds: #status experimental

F:555,562,565/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 1; Length 566;

Best Local Similarity 100.0%; Pred. No. 1,4e-10;

Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAAGFIENGWEGMIDGMYG 24

Db 346 GLFGAIAAGFIENGWEGMIDGMYG 368

RESULT 14

HM1IVH

hemagglutinin precursor - influenza A virus (strain A/Aichi/2/68)

N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: Influenza A virus

C:Date: 19-May-1994 #sequence_revision 19-May-1994 #text_change 16-Jul-1999

C:Accession: A93231; A04051

R:Verhoeven, M.; Pang, R.; Min Jou, W.; Devos, R.; Huylebroeck, D.; Saman, E.; Piers, W.

Nature 286, 771-776, 1980

A:Title: Antigenic drift between the haemagglutinin of the Hong Kong influenza strains A

A:Reference number: A93231; MUID:80254693; PMID:7402351

A:Accession: A93231

A:Molecule type: genomic RNA

A:Residues: 1-566 <VER>

A:Cross-references: GB:J02090; NID:g324131; PIDN:AA43178.1; PID:g324132

C:Superfamily: influenza virus hemagglutinin

C:Keywords: hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-16/DNA: signal sequence #status predicted <SIG>

F:17-344/Product: hemagglutinin HA1 #status predicted <HA1>

F:346-566/Product: hemagglutinin HA2 #status predicted <HA2>

F:555,562,565/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 1; Length 566;

Best Local Similarity 100.0%; Pred. No. 1,4e-10;

Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAAGFIENGWEGMIDGMYG 24

Db 346 GLFGAIAAGFIENGWEGMIDGMYG 368

RESULT 15

HM1VHM
hemagglutinin precursor - influenza A virus (strain A/Mem/102/72)
N:Contains: hemagglutinin HA1; hemagglutinin HA2
C:Species: influenza A virus
C>Date: 19-May-1994 #sequence_revision 19-May-1994 #text_change 31-Mar-2000
C:Accession: A94441; A04051
R:Seigh, M.J.; Both, G.W.; Brownlee, G.G.; Bender, V.J.; Moss, B.A.
in Structure and Variation in Influenza Virus, Laver, G., and Air, G., eds., pp.69-79, E
A:Title: The haemagglutinin gene of influenza A virus: nucleotide sequence analysis of c
A:Reference number: A94441
A:Accession: A94441
A:Molecule type: genomic RNA
A:Residues: 1-566 <SLR>
C:Superfamily: influenza virus hemagglutinin
C:Keywords: hemagglutinin; homotrimer; lipoprotein; thiolester bond
F:1-16/Domain: signal sequence #status predicted <SIG>
F:17-344/Product: hemagglutinin HA1 #status predicted <HA1>
F:346-566/Product: hemagglutinin HA2 #status predicted <HA2>
F:555,562,565/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 93.7%; Score 134; DB 1; Length 566;
Best Local Similarity 100.0%; Pred. No. 1.4e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMIDGWTG 24
|||
Db 346 GLFGAIAGFIENGWEGMIDGWTG 368

Search completed: January 30, 2004, 00:26:21
Job time : 10.2535 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 29, 2004, 23:58:27 ; Search time 5.40845 Seconds
(without alignments)
208.681 Million cell updates/sec

Title: US-09-461-684C-3
Perfect score: 143
Sequence: 1 CGLFGALAGFIENGMEGMIDGMYG 24

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues
Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_41:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length	ID	Description
1	134	93.7	550 1 HEMA_IADH1	P12582 influenza a
2	134	93.7	550 1 HEMA_IADH2	P12583 influenza a
3	134	93.7	550 1 HEMA_IADH3	P12584 influenza a
4	134	93.7	550 1 HEMA_IADH4	P12585 influenza a
5	134	93.7	550 1 HEMA_IADH5	P12586 influenza a
6	134	93.7	550 1 HEMA_IADH6	P12587 influenza a
7	134	93.7	550 1 HEMA_IADH7	P12588 influenza a
8	134	93.7	550 1 HEMA_IADH8	P43257 influenza a
9	134	93.7	550 1 HEMA_IADH9	P43258 influenza a
10	134	93.7	550 1 HEMA_IADH10	P43260 influenza a
11	134	93.7	550 1 HEMA_IADH11	P11133 influenza a
12	134	93.7	550 1 HEMA_IADH12	P11134 influenza a
13	134	93.7	556 1 HEMA_IADH13	P03437 influenza a
14	134	93.7	556 1 HEMA_IADH14	P03442 influenza a
15	134	93.7	556 1 HEMA_IADH15	P26134 influenza a
16	134	93.7	556 1 HEMA_IADH16	P26138 influenza a
17	134	93.7	556 1 HEMA_IADH17	P03449 influenza a
18	134	93.7	556 1 HEMA_IADH18	P03439 influenza a
19	134	93.7	556 1 HEMA_IADH19	P03436 influenza a
20	133	93.0	561 1 HEMA_IADH20	P12581 influenza a
21	133	93.0	561 1 HEMA_IADH21	P12439 influenza a
22	132	92.3	565 1 HEMA_IADH22	P17000 influenza a
23	132	92.3	565 1 HEMA_IADH23	P17002 influenza a
24	132	92.3	566 1 HEMA_IADH24	P03440 influenza a
25	132	92.3	566 1 HEMA_IADH25	P03439 influenza a
26	132	92.3	567 1 HEMA_IADH26	P03435 influenza a
27	131	91.6	550 1 HEMA_IADH27	P12589 influenza a
28	131	91.6	560 1 HEMA_IADH28	P03458 influenza a
29	131	91.6	565 1 HEMA_IADH29	P16994 influenza a
30	131	91.6	565 1 HEMA_IADH30	P16995 influenza a
31	131	91.6	565 1 HEMA_IADH31	P16996 influenza a
32	131	91.6	565 1 HEMA_IADH32	P16996 influenza a
33	131	91.6	565 1 HEMA_IADH33	P16558 influenza a

34	131	91.6	565 1 HEMA_IADH34	P16997 influenza a
35	131	91.6	565 1 HEMA_IADH35	P16998 influenza a
36	131	91.6	565 1 HEMA_IADH36	P16999 influenza a
37	131	91.6	565 1 HEMA_IADH37	P08011 influenza a
38	131	91.6	565 1 HEMA_IADH38	P17001 influenza a
39	131	91.6	566 1 HEMA_IADH39	P26141 influenza a
40	131	91.6	570 1 HEMA_IADH40	P26094 influenza a
41	131	91.6	570 1 HEMA_IADH41	P26095 influenza a
42	131	91.6	570 1 HEMA_IADH42	P26096 influenza a
43	131	91.6	570 1 HEMA_IADH43	P26097 influenza a
44	131	91.6	570 1 HEMA_IADH44	P26098 influenza a
45	131	91.6	570 1 HEMA_IADH45	P26099 influenza a

ALIGNMENTS

RESULT 1
HEMA_IADH1 STANDARD; PRT; 550 AA.
AC P12582; Q84021; Q84022;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-OCT-1989 (Rel. 12, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
DE Hemagglutinin HA2 chain] (Fragment).
GN HA.
OS Influenza A virus (strain A/Duck/Hokkaido/5/77).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCHI_TaxID=11357;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87265458; PubMed=2440178;
RA Kida H., Kawoka Y., Nave C.W., Webster R.G.;
RT "Antigenic and genetic conservation of H3 influenza virus in wild
RT ducks";
RL Virology 159:109-119 (1987).
CC -!- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC -!- CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -!- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -!- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
CC
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CC
CC EMBL; M16737; AAA43143.1; -.
DR HSSP; P03437; 3HWG.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutn; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Hemagglutinin; Glycoprotein.
FT NON TER 1
FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CHAIN 330 550 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 550 AA; 61705 MW; 767ACEF716FC969A CRC64;
Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.2e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAAGFIENGEGMIDGWTG 24
 DB 330 GLFGAAGFIENGEGMIDGWTG 352

RESULT 2

HEMA_IADH2 STANDARD; PRT; 550 AA.

AC P12583; Q84011; (Created)
 DT 01-OCT-1989 (Rel. 12, Last sequence update)
 DT 01-APR-1990 (Rel. 14, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
 GN Hemagglutinin HA2 chain] (Fragment).
 OS Influenza A virus (strain A/Duck/Hokkaido/8/80).
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 OC Influenza A viruses; Influenzavirus A.
 OX NCBI_TaxId=11358;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87265458; PubMed=2440178;
 RA Kida H., Kawasaka Y., Naeve C.W., Webster R.G.;
 RT "Antigenic and genetic conservation of H3 influenza virus in wild
 RT ducks";
 RL Virology 159:109-119(1987).
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION.
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.

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DR EMBL; M16738; AAA43144.1; -.
 DR PIR; B27813; HMI180.
 DR HSSP; P03437; 2V1U.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTIN12.
 DR ProDom; PD000225; Hemagglutn; 1.
 KM Envelope protein; Hemagglutinin; Glycoprotein.
 FT NON TER 1
 FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
 FT CHAIN 330 550 HEMAGGLUTININ HA2 CHAIN.
 FT CARBOHYD 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CONFLICT 137 K -> N (IN PIR DATA BANK).
 SQ SEQUENCE 550 AA; 61659 MW; A107023AC9CC353 CRC64;

Query Match 93.7%; Score 134; DB 1; Length 550;
 Best Local Similarity 100.0%; Pred. No. 1.2e-10;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAAGFIENGEGMIDGWTG 24
 DB 330 GLFGAAGFIENGEGMIDGWTG 352

RESULT 3
 HEMA_IADH3 STANDARD; PRT; 550 AA.

AC P12584; Q84012; Q89793;
 DT 01-OCT-1989 (Rel. 12, Created)
 DT 01-OCT-1989 (Rel. 12, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
 GN Hemagglutinin HA2 chain] (Fragment).
 OS Influenza A virus (strain A/Duck/Hokkaido/33/80).
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 OC Influenza A viruses; Influenzavirus A.
 OX NCBI_TaxId=11359;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87265458; PubMed=2440178;
 RA Kida H., Kawasaka Y., Naeve C.W., Webster R.G.;
 RT "Antigenic and genetic conservation of H3 influenza virus in wild
 RT ducks";
 RL Virology 159:109-119(1987).
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION.
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.

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DR EMBL; M16739; AAA43145.1; -.
 DR HSSP; P03437; 2V1U.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTIN12.
 DR ProDom; PD000225; Hemagglutn; 1.
 KM Envelope protein; Hemagglutinin; Glycoprotein.
 FT NON TER 1
 FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
 FT CHAIN 330 550 HEMAGGLUTININ HA2 CHAIN.
 FT CARBOHYD 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 550 AA; 61577 MW; 6C30BP67CFDC97DE CRC64;

Query Match 93.7%; Score 134; DB 1; Length 550;
 Best Local Similarity 100.0%; Pred. No. 1.2e-10;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAAGFIENGEGMIDGWTG 24
 DB 330 GLFGAAGFIENGEGMIDGWTG 352

RESULT 4
 HEMA_IADH4 STANDARD; PRT; 550 AA.

AC P12585; Q84013; Q84014;
 DT 01-OCT-1989 (Rel. 12, Created)
 DT 01-OCT-1989 (Rel. 12, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
 GN Hemagglutinin HA2 chain] (Fragment).
 OS Influenza A virus (strain A/Duck/Hokkaido/7/82).
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 OC Influenza A viruses; Influenzavirus A.
 OX NCBI_TaxId=11360;

```

RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87265458; PubMed=2440178;
RA Kida H., Kawaka Y., Naewe C.W., Webster R.G.;
RT Antigenic and genetic conservation of H3 influenza virus in wild
RT ducks.
RL Virology 159:109-119(1987).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; M16740; AAA43146.1; -.
DR HSSP; P03437; 2VIU.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTIN2.
DR ProDom; PD000225; Hemagglutn; 1.
DR Envelope protein; Hemagglutinin; Glycoprotein.
KM NON TER 1 1
FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CARBOHYD 330 550 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 8 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 550 AA; 61664 MW; A16B2C8CBBD9D0 CRC64;

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.2e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMTDGMWG 24
DB 330 GLFGAIAGFIENGWEGMTDGMWG 352

RESULT 5
HEMA_IADH5 STANDARD; PRT; 550 AA.
ID HEMA_IADH5 Q84015; Q84016;
AC P12586; Q84015; Q84016;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-APR-1990 (Rel. 14, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
DE Hemagglutinin HA2 chain] (Fragment).
GN HA.
OS Influenza A virus (strain A/Duck/Hokkaido/21/82).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_Taxid=11361;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87265458; PubMed=2440178;
RA Kida H., Kawaka Y., Naewe C.W., Webster R.G.;
RT Antigenic and genetic conservation of H3 influenza virus in wild
RT ducks.
RL Virology 159:109-119(1987).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.

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CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; M16741; AAA43147.1; -.
DR HSSP; P03437; 2VIU.
DR PIR; E27813; HMI21.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTIN2.
DR ProDom; PD000225; Hemagglutn; 1.
KM Envelope protein; Hemagglutinin; Glycoprotein.
FT NON TER 1 1
FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CARBOHYD 330 550 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 7 7 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 8 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 178 179 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CONFLICT 388 K -> T (IN PIR DATA BANK).
SQ SEQUENCE 550 AA; 61856 MW; 48401C867A15BFC8 CRC64;

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.2e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMTDGMWG 24
DB 330 GLFGAIAGFIENGWEGMTDGMWG 352

RESULT 6
HEMA_IADH6 STANDARD; PRT; 550 AA.
ID HEMA_IADH6 Q84017;
AC P12587; Q84017;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-APR-1990 (Rel. 14, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
DE Hemagglutinin HA2 chain] (Fragment).
GN HA.
OS Influenza A virus (strain A/Duck/Hokkaido/9/85).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_Taxid=11362;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87265458; PubMed=2440178;
RA Kida H., Kawaka Y., Naewe C.W., Webster R.G.;
RT Antigenic and genetic conservation of H3 influenza virus in wild
RT ducks.
RL Virology 159:109-119(1987).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
CC -----
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DR EMBL; M16742; AAA43148.1; -.
DR PIR; F27813; HMTV98.
DR HSBB; P03437; IHGJ.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; Hemagglutin; 1.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Hemagglutinin; Glycoprotein.
FT FT CHAIN 1 1
FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CONFLICT 8 Y -> N (IN PIR DATA BANK).
SQ SEQUENCE 550 AA; 61711 MW; 67BCDB5F44736CFE CRC64;

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.2e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DQ 2 GLFGALGFINGWEGMIDGWYG 24
|||
Db 330 GLFGALGFINGWEGMIDGWYG 352

RESULT 7
HEMA_IADH7 STANDARD: PRT; 550 AA.

ID HEMA_IADH7
AC P12588; Q84018; Q89470;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-OCT-1989 (Rel. 12, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
DE Hemagglutinin HA2 chain] (Fragment).
GN HA.
OS Influenza A virus (strain A/Duck/Hokkaido/10/85).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_taxid=11363;
RN RN
RP MEDLINE=67265458; PubMed=2440178;
RX Kida H., Kawakita Y., Naeye C.W., Webster R.G.;
RA "Antigenic and genetic conservation of H3 influenza virus in wild
RT ducks.";
RL Virology 159:109-119(1987).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: MONOMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
(HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
CC
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or send an email to license@sib-sib.ch).
CC
CC EMBL; M16743; AAA43149.1; -.
DR HSBB; P03437; JHMG.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; Hemagglutin; 1.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Hemagglutinin; Glycoprotein.

FT	NON TER	1	1		Hemagglutinin HA1 chain.
FT	CHAIN	1	328		Hemagglutinin HA2 chain.
FT	CARBOHYD	330	550		
FT	CARBOHYD	8	8		N-LINKED (GLCNAC . .) (POTENTIAL).
FT	CARBOHYD	22	22		N-LINKED (GLCNAC . .) (POTENTIAL).
FT	CARBOHYD	38	38		N-LINKED (GLCNAC . .) (POTENTIAL).
FT	CARBOHYD	165	165		N-LINKED (GLCNAC . .) (POTENTIAL).
FT	CARBOHYD	285	285		N-LINKED (GLCNAC . .) (POTENTIAL).
FT	CARBOHYD	483	483		N-LINKED (GLCNAC . .) (POTENTIAL).
QY	SEQUENCE	550 AA;	61761 MW;	6EF81793281D53EB CRC64;	(POTENTIAL).
Query Match					
Db	Best Local Similarity	93.7%;	Score 134;	DB 1;	Length 550;
	Matches 22;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
<p>2 GLFGAIGFIENGWEGMIDGMYG 24 330 GLFGAIGFIENGWEGMIDGMYG 352</p>					
RESULT 8					
ID	HEMA_IADHK	STANDARD;	PRT;	550 AA.	
AC	P43257;				
DT	01-NOV-1995 (Rel. 32, Created)				
DT	01-NOV-1995 (Rel. 32, Last sequence update)				
DT	16-OCT-2001 (Rel. 40, Last annotation update)				
Dc	Hemagglutinin Precursor [Contains: Hemagglutinin HA1 chain; Hemagglutinin HA2 chain] (Fragment).				
De	HA.				
GN	Influenza A virus (strain A/Duck/Hong Kong/7/75).				
OC	Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;				
OC	Influenza A viruses; Influenzavirus A.				
OX	NCBI_TaxID=11364;				
RN	[1]				
RP	SEQUENCE FROM N.A.				
RA	MEDLINE=91341491; PubMed=1875195;				
RA	Yaoda U., Shortridge K.F., Shimizu Y., Kida H.;				
RT	"Molecular evidence for a role of domestic ducks in the introduction of avian H3 influenza viruses to pigs in southern China, where the A/hong Kong/68 (H3N2) strain emerged."				
RT	A/hong Kong/68 (H3N2) strain emerged."				
RL	J. Gen. Virol. 72:2007-2010(1991).				
CC	-1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO CELL RECEPTORS AND FOR INITIATING INFECTION.				
CC	-1- SUBUNIT: HOMOTIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS (HA1 AND HA2) LINKED BY A DISULFIDE BOND.				
CC	-1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.				
CC	-----				
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CC	-----				
DR	EMBL; D00929; BAA00769.1; -.				
DR	HSSP; P03437; AVIU.				
DR	InterPro; IPR001364; Hemagglutin.				
DR	Pfam; PF00509; Hemagglutinin.1.				
DR	PRINTS; PR00329; HEMAGGLUTIN12.				
DR	ProDom; PD000225; Hemagglutin.1.				
KW	Envelope protein; Hemagglutinin; Glycoprotein.				
FT	NON TER	1	1		
FT	CHAIN	1	328		Hemagglutinin HA1 chain.
FT	CHAIN	1	328		Hemagglutinin HA2 chain.
FT	CARBOHYD	330	550		
FT	CARBOHYD	8	8		N-LINKED (GLCNAC . .) (POTENTIAL).
FT	CARBOHYD	22	22		N-LINKED (GLCNAC . .) (POTENTIAL).
FT	CARBOHYD	38	38		N-LINKED (GLCNAC . .) (POTENTIAL).
FT	CARBOHYD	165	165		N-LINKED (GLCNAC . .) (POTENTIAL).
FT	CARBOHYD	285	285		N-LINKED (GLCNAC . .) (POTENTIAL).
FT	CARBOHYD	483	483		N-LINKED (GLCNAC . .) (POTENTIAL).
QY	SEQUENCE	550 AA;	61549 MW;	864639B829F51B99 CRC64;	(POTENTIAL).

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.2e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAGFIENGWEGMIDGWTG 24
Db 330 GLFGAIAGFIENGWEGMIDGWTG 352

RESULT 9

HEMA_IADHL STANDARD; PRT; 550 AA.
AC P43258;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
Hemagglutinin HA2 chain] (Fragment).
GN HA.
OS Influenza A virus (strain A/Duck/Hong Kong/54/76).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=45412;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=91341491; PubMed=1875195;
RA Yasuda J., Shortridge K.F., Shimizu Y., Kida H.;
RT "Molecular evidence for a role of domestic ducks in the introduction
of avian H3 influenza viruses to pigs in southern China, where the
A/Hong Kong/68 (H3N2) strain emerged.";
RT J. Gen. Virol. 72:2007-2010(1991)
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
(HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.

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CC EMBL; D00931; BAA00771.1; -.
DR HSSP; P03437; 2VIT.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Hemagglutinin; Glycoprotein.
FT CHAIN 1 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CARBOHYD 8 550 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 22 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 550 AA; 61718 MW; A351C56789E4B59A CRC64;

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.2e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAGFIENGWEGMIDGWTG 24
Db 330 GLFGAIAGFIENGWEGMIDGWTG 352

RESULT 10

HEMA_IAGHK STANDARD; PRT; 550 AA.
AG P43260;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
Hemagglutinin HA2 chain] (Fragment).
GN HA.
OS Influenza A virus (strain A/Goose/Hong Kong/10/76).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=45414;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=91341491; PubMed=1875195;
RA Yasuda J., Shortridge K.F., Shimizu Y., Kida H.;
RT "Molecular evidence for a role of domestic ducks in the introduction
of avian H3 influenza viruses to pigs in southern China, where the
A/Hong Kong/68 (H3N2) strain emerged.";
RT J. Gen. Virol. 72:2007-2010(1991).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
(HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.

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CC EMBL; D00930; BAA00770.1; -.
DR HSSP; P03437; 2VIT.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Hemagglutinin; Glycoprotein.
FT CHAIN 1 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CARBOHYD 8 550 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 22 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 550 AA; 61676 MW; 9A1B094DA28BACD2 CRC64;

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 1.2e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAGFIENGWEGMIDGWTG 24
Db 330 GLFGAIAGFIENGWEGMIDGWTG 352

RESULT 11

HEMA_IAGHZ STANDARD; PRT; 550 AA.
AC P11133; Q84019; Q84020;
DT 01-JUL-1989 (Rel. 11, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin [Contains: Hemagglutinin HA1 chain; Hemagglutinin HA2
chain] (Fragment).
GN HA.

```

OS Influenza A virus (strain A/Swine/Hong Kong/81/78).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxId=11497;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=88101364; PubMed=3336940;
RA Kida H., Shortridge K.F., Webster R.G.;
RT "Origin of the hemagglutinin gene of H3N2 influenza viruses from pigs
in China.";
RL Virology 162:160-166(1988).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
(HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
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CC -----
CC EMBL; M19057; AAA43212.1; -.
CC HSSP; P03437; 2V1U.
CC InterPro; IPR001364; Hemagglutn.
CC Pfam; PF00509; Hemagglutinin; 1.
CC PRINTS; PR00329; HEMAGGLUTN12.
CC ProDom; PD000225; Hemagglutn; 1.
CC Hemagglutinin; Envelope protein; Glycoprotein.
KW NON TER
FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CHAIN 1 550 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 8 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 550 AA; 61437 MW; 1F2A7E758C531CE8 CRC64;

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. NO. 1.2e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLEGAIAFIENGWEGMIDGWYG 24
DB 330 GLEGAIAFIENGWEGMIDGWYG 352

RESULT 12
HEMA_IJAZH3 STANDARD; PRT; 550 AA.
AC P1134; Q84025; Q84026;
DT 01-JUL-1989 (Rel. 11, Created)
DT 01-JUL-1989 (Rel. 11, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin [Contains: Hemagglutinin HA1 chain; Hemagglutinin HA2
chain] (Fragment).
GN HA.
OS Influenza A virus (strain A/Swine/Hong Kong/126/82).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxId=11498;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=88101364; PubMed=3336940;
RA Kida H., Shortridge K.F., Webster R.G.;
RT "Origin of the hemagglutinin gene of H3N2 influenza viruses from pigs
in China.";
RL Virology 162:160-166(1988).

```

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CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
(HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
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or send an email to license@isb-sib.ch).
CC -----
CC EMBL; M19056; AAA43211.1; ALT_TERM.
CC HSSP; P03437; 2V1U.
CC InterPro; IPR001364; Hemagglutn.
CC Pfam; PF00509; Hemagglutinin; 1.
CC PRINTS; PR00329; HEMAGGLUTN12.
CC ProDom; PD000225; Hemagglutn; 1.
CC Hemagglutinin; Envelope protein; Glycoprotein.
KW NON TER
FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CHAIN 1 550 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 8 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 550 AA; 61580 MW; 991F6D8BC02F24F2 CRC64;

Query Match 93.7%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. NO. 1.2e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLEGAIAFIENGWEGMIDGWYG 24
DB 330 GLEGAIAFIENGWEGMIDGWYG 352

RESULT 13
HEMA_IAAIC STANDARD; PRT; 566 AA.
AC P03437;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
Hemagglutinin HA2 chain].
GN HA.
OS Influenza A virus (strain A/Aichi/2/68).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxId=150147;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=80254693; PubMed=7402351;
RA Verhoeven M., Fang R., Min Jou W., Devos R., Huylbroeck D.,
Saman B., Fiers W.;
RT "Antigenic drift between the haemagglutinin of the Hong Kong
influenza strains A/Aichi/2/68 and A/Victoria/3/75.";
RL Nature 286:771-776(1980).
RN [2]
RP X-RAY CRYSTALLOGRAPHY (3.0 ANGSTROMS).
RX MEDLINE=81123029; PubMed=7464906;
RA Wilson I.A., Skehel J.J., Wiley D.C.;
RT "Structure of the haemagglutinin membrane glycoprotein of influenza
virus at 3-A resolution.";
RL Nature 289:366-373(1981).
RN [3]
RP X-RAY CRYSTALLOGRAPHY.
RX MEDLINE=88232903; PubMed=3374584;

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RA  Weis W.I., Brown J.H., Cusack S.C., Paulson J.C., Skehel J.J.,
RT  "Structure of the influenza virus haemagglutinin complexed with its
RL  receptor, sialic acid."
RL  Nature 333:426-431(1988).
RN  [4]
RP  X-RAY CRYSTALLOGRAPHY OF A MUTANT WITH GLY-457.
RX  MEDLINE=90107940; Pubmed=2295311;
RA  Weis W.I., Cusack S.C., Brown J.H., Daniels R.S., Skehel J.J.,
RT  "The structure of a membrane fusion mutant of the influenza virus
RL  haemagglutinin."
RL  EMBO J. 9:17-24(1990).
RN  [5]
RP  X-RAY CRYSTALLOGRAPHY.
RX  MEDLINE=90230310; Pubmed=2329580;
RA  Weis W.I., Bruegger A.T., Skehel J.J., Wiley D.C.;
RT  "Refinement of the influenza virus haemagglutinin by simulated
RL  annealing."
RL  J. Mol. Biol. 212:737-761(1990).
RN  [6]
RP  X-RAY CRYSTALLOGRAPHY (2.5 ANGSTROMS).
RX  MEDLINE=94352388; Pubmed=8072525;
RA  Bullough P.A., Hughson F.M., Skehel J.J., Wiley D.C.;
RT  "Structure of influenza haemagglutinin at the pH of membrane fusion."
RL  Nature 371:37-43(1994).
RN  [7]
RP  X-RAY CRYSTALLOGRAPHY (3.25 ANGSTROMS).
RX  MEDLINE=98120975; Pubmed=9461077;
RA  Fleury D., Wharton S.A., Skehel J.J., Knossow M., Bizebard T.;
RT  "Antigen distortion allows influenza virus to escape neutralization."
RL  Nat. Struct. Biol. 5:119-123(1998).
CC  -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC  -1- CELL RECEPTORS AND FOR INITIATING INFECTION.
CC  -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC  (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC  -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
CC  -----
CC  This SWISS-PROT entry is copyright. It is produced through a collaboration
CC  between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC  the European Bioinformatics Institute. There are no restrictions on its
CC  use by non-profit institutions as long as its content is in no way
CC  modified and this statement is not removed. Usage by and for commercial
CC  entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC  or send an email to license@isb-sib.ch).
CC  -----
DR  EMBL; J02090; AAA43178.1; -.
DR  EMBL; V01085; CAA24269.1; -.
DR  PIR; A93231; HMTVHA.
DR  PDB; 2HMG; 31-OCT-93.
DR  PDB; 3HMG; 31-OCT-93.
DR  PDB; 4HMG; 31-OCT-93.
DR  PDB; 5HMG; 31-JAN-94.
DR  PDB; 1HGD; 31-JAN-94.
DR  PDB; 1HGE; 31-JAN-94.
DR  PDB; 1HGF; 31-JAN-94.
DR  PDB; 1HGH; 31-JAN-94.
DR  PDB; 1HGI; 31-JAN-94.
DR  PDB; 1HGU; 31-JAN-94.
DR  PDB; 1HTM; 14-FEB-95.
DR  PDB; 2VIR; 29-APR-98.
DR  PDB; 2VIS; 29-APR-98.
DR  PDB; 2VIT; 29-APR-98.
DR  PDB; 2VIT; 29-APR-98.
DR  PDB; 1E08; 29-NOV-00.
DR  PDB; 1HA0; 22-DEC-99.
DR  PDB; 1J8H; 13-MAR-02.
DR  PDB; 1KEN; 24-APR-02.
DR  PDB; 1QFU; 29-DEC-99.
DR  PDB; 1QU1; 05-JAN-00.
DR  InterPro; IPR001364; Hemagglutn.
DR  Pfam; PF00509; Hemagglutinin; 1.

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DR	PRINTS; PRO00329; HEMAGGLUTIN12.		
DR	ProDom; PD000225; Hemagglutin; 1.		
KW	Envelope protein; Hemagglutinin; Glycoprotein; Signal; 3D-structure		
FT	FT SGNL	1	16
FT	FT CHAIN	17	344
FT	FT CHAIN	346	566
FT	FT DISULFID	30	482
FT	FT DISULFID	68	293
FT	FT DISULFID	80	92
FT	FT DISULFID	113	155
FT	FT DISULFID	297	321
FT	FT DISULFID	489	493
FT	FT CARBOHYD	24	24
FT	FT CARBOHYD	38	38
FT	FT CARBOHYD	54	54
FT	FT CARBOHYD	97	97
FT	FT CARBOHYD	181	181
FT	FT CARBOHYD	301	301
FT	FT CARBOHYD	499	499
FT	FT SGNL	27	35
FT	FT STRAND	40	42
FT	FT STRAND	50	52
FT	FT STRAND	55	57
FT	FT STRAND	59	60
FT	FT STRAND	67	70
FT	FT STRAND	74	76
FT	FT TURN	78	79
FT	FT HELIX	82	87
FT	FT TURN	88	88
FT	FT HELIX	90	95
FT	FT TURN	96	97
FT	FT STRAND	99	99
FT	FT STRAND	102	105
FT	FT TURN	107	108
FT	FT STRAND	116	117
FT	FT TURN	119	120
FT	FT HELIX	121	131
FT	FT STRAND	133	133
FT	FT STRAND	136	138
FT	FT TURN	144	145
FT	FT STRAND	146	147
FT	FT STRAND	152	157
FT	FT TURN	158	159
FT	FT STRAND	160	162
FT	FT TURN	165	166
FT	FT STRAND	167	169
FT	FT STRAND	171	173
FT	FT TURN	174	175
FT	FT STRAND	176	176
FT	FT STRAND	180	185
FT	FT STRAND	192	200
FT	FT HELIX	204	211
FT	FT STRAND	217	221
FT	FT STRAND	226	229
FT	FT TURN	239	239
FT	FT STRAND	240	241
FT	FT STRAND	242	242
FT	FT STRAND	245	253
FT	FT TURN	255	256
FT	FT STRAND	258	265
FT	FT STRAND	267	270
FT	FT STRAND	272	275
FT	FT STRAND	282	285
FT	FT STRAND	290	294
FT	FT STRAND	297	299
FT	FT TURN	300	301
FT	FT STRAND	302	304
FT	FT STRAND	310	311
FT	FT STRAND	318	320
FT	FT STRAND	323	324
FT	FT STRAND	331	333
FT	FT STRAND	337	337
FT	FT TURN	347	348

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FT TURN 350 350
FT STRAND 351 351
FT TURN 352 354
FT STRAND 355 355
FT STRAND 359 355
FT TURN 361 361
FT STRAND 367 373
FT TURN 374 375
FT STRAND 376 382
FT TURN 383 400
FT TURN 401 401
FT STRAND 406 407
FT HELIX 421 471
FT STRAND 472 474
FT STRAND 475 477
FT STRAND 482 485
FT HELIX 491 498
FT TURN 499 500
FT HELIX 504 515
FT TURN 518 519
SQ SEQUENCE 566 AA; 63415 MW; E395659C23CAFECA CRC64;

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Query Match 93.7%; Score 134; DB 1; Length 566;
 Best Local Similarity 100.0%; Pred. No. 1.2e-10;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 2 GLFGAAGFIENGWEGMIDGWYG 24
DB 346 GLFGAAGFIENGWEGMIDGWYG 368

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RESULT 14

HEMA_IAD3 STANDARD; PRT; 566 AA.

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AC P26134;
DT 01-MAY-1992 (Rel. 22, Created)
DT 01-MAY-1992 (Rel. 22, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HAI chain;
  Hemagglutinin HAZ chain].
GN HA.
OS Influenza A virus (strain A/Duck/Alberta/78/76).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_Taxid=11348;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=92114135; PubMed=1731092;
RA Bean W.J., Scheil M., Katz J., Kawoka Y., Naeve C., Gorman O.,
  Webster R.G.;
RT "Evolution of the H3 influenza virus hemagglutinin from human and
  nonhuman hosts.";
RL J. Virol. 66:1129-1138(1992).

```

```

CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
  CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
  (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.

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  or send an email to license@sib-sib.ch).

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CC EMBL: M73771; -; NOT_ANNOTATED_CDS.
DR HSSP; P03437; 2VIT.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.

```

```

KM Envelope protein; Hemagglutinin; Glycoprotein; Signal.
FT SIGNAL 1 16
FT CHAIN 17 344
FT CHAIN 346 566
FT CARBOHYD 23 23
FT CARBOHYD 24 24
FT CARBOHYD 38 38
FT CARBOHYD 54 54
FT CARBOHYD 181 181
FT CARBOHYD 301 301
FT CARBOHYD 499 499
SQ SEQUENCE 566 AA; 63534 MW; FE19AB6FP9415B89 CRC64;

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Query Match 93.7%; Score 134; DB 1; Length 566;
 Best Local Similarity 100.0%; Pred. No. 1.2e-10;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 2 GLFGAAGFIENGWEGMIDGWYG 24
DB 346 GLFGAAGFIENGWEGMIDGWYG 368

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RESULT 15

HEMA_IAD3 STANDARD; PRT; 566 AA.

```

AC P03442;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HAI chain;
  Hemagglutinin HAZ chain].
GN HA.
OS Influenza A virus (strain A/Duck/Ukraine/1/63).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_Taxid=11374;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=82025542; PubMed=6169439;
RA Pang R., Min Jou W., Huybrecock D., Devos R., Fiers W.;

```

```

RT "Complete structure of A/duck/Ukraine/63 influenza hemagglutinin
  gene: animal virus as progenitor of human H3 Hong Kong 1968 influenza
  hemagglutinin.";
RL Cell 25:315-323(1981).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
  CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
  (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.

```

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  or send an email to license@sib-sib.ch).

```

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CC EMBL; V01087; CAA24271.1; -.
DR PDB; 1IBN; 08-AUG-01.
DR PDB; 1IBO; 08-AUG-01.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Hemagglutinin; Glycoprotein; Signal; 3D-structure.

```

```

FT SIGNAL 1 16
FT CHAIN 17 344
FT CHAIN 346 566
FT CARBOHYD 23 24
FT CARBOHYD 38 38
FT CARBOHYD 54 54
FT CARBOHYD 97 97

```


FT CARBOHYD 181 181 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 301 301 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 499 499 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 566 AA; 63530 MW; E70F87F0AE1178F4 CRC64;

Query Match 93.7%; Score 134; DB 1; Length 566;
 Best Local Similarity 100.0%; Pred. No. 1.2e-10;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAIGFIENGWEGMIDGWTG 24
 |||||
 DB 346 GLFGAIGFIENGWEGMIDGWTG 368

Search completed: January 30, 2004, 00:20:45
 Job time : 5.40845 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 29, 2004, 23:58:52 ; Search time 24.7887 Seconds
(without alignments)
249.842 Million cell updates/sec

Title: US-09-461-684C-3
Perfect score: 143
Sequence: 1 GLEFGAIGFIENGWEGMIDGWYG 24

Scoring table: BIOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

SPTREMBL_23:*

- 1: sp_archaea:*
- 2: sp_bacteria:*
- 3: sp_fungi:*
- 4: sp_human:*
- 5: sp_invertebrate:*
- 6: sp_mammal:*
- 7: sp_mhc:*
- 8: sp_organelle:*
- 9: sp_phage:*
- 10: sp_plant:*
- 11: sp_rodent:*
- 12: sp_virus:*
- 13: sp_vertebrate:*
- 14: sp_unclassified:*
- 15: sp_virus:*
- 16: sp_bacteriophage:*
- 17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	134	93.7	384	12	Q8JK63 influenza a
2	134	93.7	566	12	Q98052 influenza a
3	134	93.7	566	12	Q8U251 influenza a
4	134	93.7	566	12	Q8OLN8 influenza a
5	134	93.7	566	12	Q67132 influenza a
6	134	93.7	566	12	Q67125 influenza a
7	134	93.7	566	12	Q8UXK3 influenza a
8	134	93.7	566	12	Q91M77 influenza a
9	134	93.7	566	12	Q9DHG0 influenza a
10	134	93.7	566	12	Q910M5 influenza a
11	134	93.7	566	12	Q67126 influenza a
12	133	93.0	301	12	Q9DXE3 influenza a
13	132	92.3	550	12	Q82498 influenza a
14	132	92.3	550	12	Q82498 influenza a
15	132	92.3	550	12	Q82753 influenza a
16	132	92.3	566	12	Q82496 influenza a

17	132	92.3	571	12	Q03909 influenza a
18	131	91.6	109	12	Q67050 influenza a
19	131	91.6	109	12	Q67053 influenza a
20	131	91.6	109	12	Q67051 influenza a
21	131	91.6	109	12	Q67052 influenza a
22	131	91.6	362	12	Q9QKD3 influenza a
23	131	91.6	362	12	Q9QKD1 influenza a
24	131	91.6	362	12	Q82513 influenza a
25	131	91.6	362	12	Q9QKD2 influenza a
26	131	91.6	362	12	Q84174 influenza a
27	131	91.6	362	12	Q82517 influenza a
28	131	91.6	365	12	Q9DL25 influenza a
29	131	91.6	367	12	Q9DL22 influenza a
30	131	91.6	368	12	Q9DL29 influenza a
31	131	91.6	369	12	Q9DL26 influenza a
32	131	91.6	369	12	P87689 influenza a
33	131	91.6	369	12	Q9DL06 influenza a
34	131	91.6	371	12	Q9DL24 influenza a
35	131	91.6	371	12	P87685 influenza a
36	131	91.6	373	12	Q9DL20 influenza a
37	131	91.6	374	12	Q9DL21 influenza a
38	131	91.6	375	12	Q9DL27 influenza a
39	131	91.6	375	12	Q9DL05 influenza a
40	131	91.6	376	12	Q9DL30 influenza a
41	131	91.6	376	12	Q9DL04 influenza a
42	131	91.6	377	12	Q9E7P6 influenza a
43	131	91.6	382	12	Q9DL03 influenza a
44	131	91.6	408	12	Q9E7P5 influenza a
45	131	91.6	409	12	Q9Q0L5 influenza a

ALIGNMENTS

RESULT 1

Q8JK63 PRELIMINARY; PRT; 384 AA.
ID Q8JK63;
AC Q8JK63;
DT 01-OCT-2002 (TREMBLrel. 22, Created)
DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Hemagglutinin (Fragment).
GN H3HA.
OS Influenza A virus (A/real/Germany/wv201r/01).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=205472;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A/real/Germany/wv01r/01;
RA Werner O., Starick B., Mueller T., Muehle R.;
RT "Characterisation of avian influenza virus isolates from wild birds from Germany.";
RT Submitted (Aug-2002) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -!- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL; AJ506781; CAD44999.1;-.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
FT NON TER 384
SQ SEQUENCE 384 AA; 42076 MW; 459731795CA5C838 CRC64;

Query Match 93.7%; Score 134; DB 12; Length 384;
Best local Similarity 100.0%; Pred. No. 5.9e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 GLEFGAIGFIENGWEGMIDGWYG 24

Db 346 GLFGAIAGFIENGWEGMIDGWYG 368

RESULT 2

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Q98052 ID Q98052 PRELIMINARY; PRT; 566 AA.
AC Q98052;
DT 01-FEB-1997 (TRENBLREL. 02, Created)
DT 01-FEB-1997 (TRENBLREL. 02, Last sequence update)
DT 01-OCT-2002 (TRENBLREL. 22, Last annotation update)
DE Hemagglutinin precursor (Fragment).
OS Influenzavirus A.
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses.
OX NCB1_Taxid=197911;
RN (1)
RP SEQUENCE FROM N.A.
RX MEDLINE=81053698; PubMed=6253883;
RA Both G.W., Sleight M.J.;
RT "Complete nucleotide sequence of the haemagglutinin gene from a human
RT Influenza virus of the Hong Kong subtype.";
RL Nucleic Acids Res. 8:2561-2575(1980).
RN (2)
RP SEQUENCE OF 17-344 FROM N.A.
RX MEDLINE=81194918; PubMed=6164798;
RA Sleight M.J., Both G.W., Underwood P.A., Bender V.J.;
RT "Antigenic drift in the hemagglutinin of the Hong Kong influenza
RT subtype: Correlation of amino acid changes with alterations in viral
RT antigenicity.";
RL J. Virol. 37:845-853(1981).
RN (3)
RP SEQUENCE OF 17-566 FROM N.A.
RX MEDLINE=82033276; PubMed=6169843;
RA Both G.W., Sleight M.J.;
RT "Conservation and variation in the hemagglutinins of Hong Kong subtype
RT influenza viruses during antigenic drift.";
RL J. Virol. 39:845-853(1981).
RN (4)
RP FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
RN CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL: J02135; AAA43189.1; -.
DR HSPF: P03437; IHGE.
DR InterPro: IPR001364; Hemagglutn.
DR Pfam: PF00509; Hemagglutinin; 1.
DR PRINTS: PR00329; HEMAGGLUTN12.
DR ProDom: PD000225; Hemagglutn; 1.
KW Envelope protein; Glycoprotein; Hemagglutinin; Signal.
FT SIGNAL 1 16
FT CHAIN 17 344
FT CHAIN 346 566
FT CHAIN 566 566
FT CHAIN 566 566
SQ SEQUENCE 566 AA; 63414 MW; C447FD465BE4FCF9 CRC64;

Query Match 93.7%; Score 134; DB 12; Length 566;
Best Local Similarity 100.0%; Pred. No. 8.8e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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OS Influenza A virus (A/pet bird/Hong Kong/1559/99 (H3N8)).
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 OC Influenza A viruses; Influenzavirus A.
 OX NCB1_Taxid=183665;

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(1)
RP SEQUENCE FROM N.A.
RC STRAIN=A/pet bird/Hong Kong/1559/99;
RA Chin P., Shortridge K.F.;
RT "Characterisation of avian H3 influenza viruses.";
RL Submitted (JUN-2002) to the EMBL/Genbank/DBJ databases.
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL: AJ427304; CAD20336.1; -.
DR InterPro: IPR001364; Hemagglutn.
DR Pfam: PF00509; Hemagglutinin; 1.
DR PRINTS: PR00329; HEMAGGLUTN12.
DR ProDom: PD000225; Hemagglutn; 1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
SQ SEQUENCE 566 AA; 63403 MW; F11C91B6A0183484 CRC64;

Query Match 93.7%; Score 134; DB 12; Length 566;
Best Local Similarity 100.0%; Pred. No. 8.8e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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RESULT 4

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Q980L8 ID Q980L8 PRELIMINARY; PRT; 566 AA.
AC Q980L8;
DT 01-JUN-2002 (TRENBLREL. 21, Created)
DT 01-JUN-2002 (TRENBLREL. 21, Last sequence update)
DT 01-MAR-2003 (TRENBLREL. 23, Last annotation update)
DE Haemagglutinin.
RN HA.
OS Influenza A virus (A/aquatic bird/Hong Kong/399/99 (H3N8)).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCB1_Taxid=183664;
RN (1)
RP SEQUENCE FROM N.A.
RC STRAIN=A/aquatic bird/Hong Kong/399/99;
RA Chin P., Shortridge K.F.;
RT "Characterisation of influenza viruses from wild aquatic birds.";
RL Submitted (JUN-2002) to the EMBL/Genbank/DBJ databases.
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL: AJ427297; CAD20332.1; -.
DR InterPro: IPR001364; Hemagglutn.
DR Pfam: PF00509; Hemagglutinin; 1.
DR ProDom: PD000225; Hemagglutn; 1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
SQ SEQUENCE 566 AA; 63412 MW; 68913C222C97B92E CRC64;

Query Match 93.7%; Score 134; DB 12; Length 566;
Best Local Similarity 100.0%; Pred. No. 8.8e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Q980L8 ID Q980L8 PRELIMINARY; PRT; 566 AA.
AC Q980L8;
DT 01-MAR-2002 (TRENBLREL. 20, Created)
DT 01-MAR-2002 (TRENBLREL. 20, Last sequence update)
DT 01-OCT-2002 (TRENBLREL. 22, Last annotation update)
DE Haemagglutinin.
RN HA.

Query Match 93.7%; Score 134; DB 12; Length 566;
Best Local Similarity 100.0%; Pred. No. 8.8e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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067132
ID 067132 PRELIMINARY; PRT: 566 AA.
AC 067132;
DT 01-NOV-1996 (TREMblrel. 01, Created)
DT 01-NOV-1996 (TREMblrel. 01, Last sequence update)
DT 01-OCT-2002 (TREMblrel. 22, Last annotation update)
DE Hemagglutinin.
GN HA.
OS Influenza A virus (strain A/Aichi/2/68).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=150147;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A/Aichi/2/68;
RA Min J.W., Verhoeven M., Fang R.-X., Devos R., Huybrecock D.,
RA Fiers W.;
RT "Shift and drift in influenza viruses.";
RL (in) Carlisle M.J., Collins J.F., Moseley B.E. B. (eds.);
RL SYMPOSIUM OF THE SOCIETY FOR GENERAL MICROBIOLOGY, pp.285-311,
RL Cambridge University Press, New York (1981)
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL; M55059; AA43239.1; -.
DR HSSP; P03437; IHGS.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
FT CHAIN 1 346
FT NEURAMINIDASE.
SQ SEQUENCE 566 AA; 63441 MW; E5D1B97DF96FECA CRC64;
Query Match 93.7%; Score 134; DB 12; Length 566;
Best Local Similarity 100.0%; Pred. No. 8.8e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 2 GLFGAAGFIENGWEGMIDGWYG 24
DB 346 GLFGAAGFIENGWEGMIDGWYG 368
RESULT 6
067125
ID 067125 PRELIMINARY; PRT: 566 AA.
AC 067125;
DT 01-NOV-1996 (TREMblrel. 01, Created)
DT 01-NOV-1996 (TREMblrel. 01, Last sequence update)
DT 01-OCT-2002 (TREMblrel. 22, Last annotation update)
DE Hemagglutinin.
GN HA.
OS Influenzavirus A.
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses.
OX NCBI_TaxID=197911;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A/Seal/M/3911/92;
RA MEDLINE=95146951; PubMed=784453;
RA Callan R.U., Early G., Kida H., Hinshaw V.S.;
RT "The appearance of H3 influenza viruses in seals.";
RL J. Gen. Virol. 76:199-203(1995).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL; L31949; AAA64229.1; -.
DR HSSP; P03437; 2VU.

DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
SQ SEQUENCE 566 AA; 63456 MW; AE556302A9EBB99F CRC64;
Query Match 93.7%; Score 134; DB 12; Length 566;
Best Local Similarity 100.0%; Pred. No. 8.8e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 2 GLFGAAGFIENGWEGMIDGWYG 24
DB 346 GLFGAAGFIENGWEGMIDGWYG 368
RESULT 7
08UXR3
ID 08UXR3 PRELIMINARY; PRT: 566 AA.
AC 08UXR3;
DT 01-MAR-2002 (TREMblrel. 20, Created)
DT 01-MAR-2002 (TREMblrel. 20, Last sequence update)
DT 01-OCT-2002 (TREMblrel. 22, Last annotation update)
DE Hemagglutinin.
GN HA.
OS Influenza A virus (A/swine/Potsdam/35/82 (H3N2)).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=183769;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A/swine/Potsdam/35/82;
RA Groetzingen I., Sues J., Groetzingen C.;
RT "Evolution of european human and porcine influenza viruses.";
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL; AJ252132; CAC81018.1; -.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
SQ SEQUENCE 566 AA; 63529 MW; 6AA44C84B4DDE68A CRC64;
Query Match 93.7%; Score 134; DB 12; Length 566;
Best Local Similarity 100.0%; Pred. No. 8.8e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 2 GLFGAAGFIENGWEGMIDGWYG 24
DB 346 GLFGAAGFIENGWEGMIDGWYG 368
RESULT 8
091MA7
ID 091MA7 PRELIMINARY; PRT: 566 AA.
AC 091MA7;
DT 01-DEC-2001 (TREMblrel. 19, Created)
DT 01-DEC-2001 (TREMblrel. 19, Last sequence update)
DT 01-MAR-2003 (TREMblrel. 23, Last annotation update)
DE Hemagglutinin.
OS Influenza A virus (A/Hong Kong/1/68 (H3N2)).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=108859;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A/Hong Kong/1/68 (H3N2);
RA MEDLINE=21287244; PubMed=11371620;

RA	Brown E.G., Liu H., Kit L.C., Baird S., Nesraliah M.;
RT	"Pattern of mutation in the genome of influenza A virus on adaptation
RT	to increased virulence in the mouse lung; Identification of functional
RT	themes";
RL	Proc. Natl. Acad. Sci. U.S.A. 98:6883-6888 (2001).
CC	-1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC	CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC	-1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC	(HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC	-1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR	EMBL; AF348176; AAKS1718.1; -.
DR	InterPro; IPR001364; Hemagglutn.
DR	Pfam; PF00509; Hemagglutinin; 1.
DR	PRINTS; PR00329; HEMAGGLUTN12.
DR	ProDom; PD000225; Hemagglutn; 1.
KW	Envelope protein; Glycoprotein; Hemagglutinin.
SQ	SEQUENCE 566 AA; 63387 MW; 01BB0D465BE158E1 CRC64;

QY	Query Match	93.7%; Score 134;	DB 12;	Length 566;
Db	Best Local Similarity	100.0%; Pred. No. 8.8e-10;		
	Matches 23; Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0

Db	346 GLFGAIAGFIENGWGMDGWG 368
QY	2 GLFGAIAGFIENGWGMDGWG 24

RESULT 9	
O9DHGO	PRELIMINARY; PRT; 566 AA.
ID O9DHGO	
AC O9DHGO	
DT 01-MAR-2001 (TREMBlrel. 16, Created)	
DT 01-MAR-2001 (TREMBlrel. 16, Last sequence update)	
DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)	
DS Haemagglutinin precursor.	
OS Influenza A virus H3N2.	
OC Viruses; sRNA negative-strand viruses; Orthomyxoviridae;	
OC Influenza A viruses; Influenzavirus A.	
OX NCBI_TaxID=41857;	
RN [1]	
RN SEQUENCE FROM N.A.	
RC STRAIN=clone 7a;	
RA Mohsin M.A., Morris S.J., Smith H., Sweet C.;	
RT "Influenza virus-induced apoptosis: a dual role for viral	
RT neuraminidase.";	
RL Submitted (JUN-2000) to the EMBL/Genbank/DBJ databases.	
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO	
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).	
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS	
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).	
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.	
DR EMBL; AJ289703; CAC18525.1; -.	
DR HSSP; P03437; 2VU.	
DR InterPro; IPR001364; Hemagglutn.	
DR Pfam; PF00509; Hemagglutinin; 1.	
DR PRINTS; PR00329; HEMAGGLUTN12.	
DR ProDom; PD000225; Hemagglutn; 1.	
KW Envelope protein; Glycoprotein; Hemagglutinin; Signal.	
FT SIGNAL 16	POTENTIAL.
SO SEQUENCE 566 AA; 63356 MW; 0BA681929300F72F CRC64;	

QY	Query Match	93.7%; Score 134;	DB 12;	Length 566;
Db	Best Local Similarity	100.0%; Pred. No. 8.8e-10;		
	Matches 23; Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0

Db	346 GLFGAIAGFIENGWGMDGWG 368
QY	2 GLFGAIAGFIENGWGMDGWG 24

RESULT 10	
O91OMS	PRELIMINARY; PRT; 566 AA.
ID Q91OMS	

0910M5 ;
01-DEC-2001 (Tremblrel. 19, Created)
01-DEC-2001 (Tremblrel. 19, Last sequence update)
01-MAR-2003 (Tremblrel. 23, Last annotation update)
Hemagglutinin.
OS Influenza A virus (A/Hong Kong/1/68 (H3N2)).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
CX NCBI_TaxID=108859;
[1]
RN SEQUENCE FROM N.A.
RP STRAIN=A/Hong Kong/1/68 (H3N2);
RX MEDLINE=21287244; PubMed=11371620;
RA Brown E.G., Liu H., Kit L.C., Baird S., Neerallah M.;
RT "Pattern of mutation in the genome of influenza A virus on adaptation
RT to increased virulence in the mouse lung: Identification of functional
RT themes.";
RL Proc. Natl. Acad. Sci. U.S.A. 98:6883-6888(2001).
RC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL; AF348179; AAK51721.1; -;
DR EMBL; AF348178; AAK51720.1; -;
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
SQ SEQUENCE 566 AA; 63530 MW; 7CB95BAP1BE6974 CRC64;

Query Match 93.7%; Score 134; DB 12; Length 566;
Best Local Similarity 100.0%; Pred. No. 8; 8e-10;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

2 GLFGAIGFTENGWEGMIDGMYG 24
Dy |||||||||||||||||||||
Db 346 GLFGAIGFTENGWEGMIDGMYG 368

RESULT 11
067126 PRELIMINARY; PRT; 566 AA.
AC 067126;
DT 01-NOV-1996 (Tremblrel. 01, Created)
DT 01-NOV-1996 (Tremblrel. 01, Last sequence update)
DT 01-OCT-2002 (Tremblrel. 22, Last annotation update)
DE Hemagglutinin.
GN HA.
OS Influenzavirus A.
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses.
CX NCBI_TaxID=197911;
RX [1]
RN SEQUENCE FROM N.A.
RP STRAIN=A/Seal/MA/3984/92;
RX MEDLINE=95146951; PubMed=7844533;
RA Callan R.J., Early G., Kida H., Hinshaw V.S.;
RT "The appearance of H3 influenza viruses in seals.";
RL J. Gen. Virol. 76:199-203(1995).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL; L32024; AAA6428.1; -;
DR HSSP; P03437; 2VIU.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.

KW Envelope protein; Glycoprotein; Hemagglutinin.
 SQ SEQUENCE 566 AA; 63441 MW; 590576CB4CEE7D08 CRC64;
 Query Match 93.7%; Score 134; DB 12; Length 566;
 Best Local Similarity 100.0%; Pred. No. 8.8e-10;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAIAFGIENGWEGMIDGWYG 24
 |||
 DB 346 GLFGAIAFGIENGWEGMIDGWYG 368

RESULT 12
 O9DXE3 PRELIMINARY; PRT; 301 AA.
 AC O9DXE3;
 DT 01-MAR-2001 (TREMBlrel. 16, Created)
 DT 01-MAR-2001 (TREMBlrel. 16, Last sequence update)
 DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
 DE Hemagglutinin (Fragment).
 GN HAI.
 OS Influenza A virus (A/Shorebird/Taiwan/31-4/99).
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 OC Influenza A viruses; Influenzavirus A.
 OX NCBI_TaxId=140665;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A/Shorebird/Taiwan/31-4/99;
 RA Lee M.S., Cheng P.C., Shien J.H., Cheng M.C., Lee L.H., Shien H.K.;
 RT "Identification and subtyping of avian influenza virus by reverse
 RT transcription-polymerase chain reaction.";
 RL Submitted (OCT-2000) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HAI AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; AF311750; AAC33016.1; -
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR Prodom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 FT NON_TER 1 301
 FT NON_TER 1 301
 SQ SEQUENCE 301 AA; 32701 MW; 62A403758B764D57 CRC64;

Query Match 93.0%; Score 133; DB 12; Length 301;
 Best Local Similarity 95.7%; Pred. No. 6.2e-10;
 Matches 22; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAIAFGIENGWEGMIDGWYG 24
 |||
 DB 250 GLFGAIAFGIENGWEGMIDGWYG 272

RESULT 13
 O82499 PRELIMINARY; PRT; 550 AA.
 AC O82499;
 DT 01-NOV-1996 (TREMBlrel. 01, Created)
 DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
 DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
 DE Hemagglutinin HAI and HA2 (Fragment).
 OS Influenzavirus A.
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 OC Influenza A viruses.
 OX NCBI_TaxId=197911;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A/Philippines/2/82/BS;
 RA Hartley C.A., Ward A.C., Anders E.M.;
 RT "Virulence of influenza virus for mice is associated with loss of

RT oligosaccharide from the hemagglutinin molecule.";
 RL Submitted (APR-1994) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HAI AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; U08859; AAA18782.1; -
 DR HSSP; P03437; 2YTU.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR Prodom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 FT NON_TER 1 550
 FT CHAIN 1 328 HAI.
 FT CHAIN 1 550 HA2.
 SQ SEQUENCE 550 AA; 61772 MW; 50BD62B6BF8E11FD8 CRC64;

Query Match 92.3%; Score 132; DB 12; Length 550;
 Best Local Similarity 95.7%; Pred. No. 1.6e-09;
 Matches 22; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAIAFGIENGWEGMIDGWYG 24
 |||
 DB 330 GLFGAIAFGIENGWEGMIDGWYG 352

RESULT 14
 O82498 PRELIMINARY; PRT; 550 AA.
 AC O82498;
 DT 01-NOV-1996 (TREMBlrel. 01, Created)
 DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
 DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
 DE Hemagglutinin HAI and HA2 (Fragment).
 OS Influenzavirus A.
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 OC Influenza A viruses.
 OX NCBI_TaxId=197911;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A/Philippines/2/82;
 RA Hartley C.A., Ward A.C., Anders E.M.;
 RT "Virulence of influenza virus for mice is associated with loss of
 RT oligosaccharide from the hemagglutinin molecule.";
 RL Submitted (APR-1994) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HAI AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; U08858; AAA18781.1; -
 DR HSSP; P03437; 2YTU.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR Prodom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 FT NON_TER 1 550
 FT CHAIN 1 328 HAI.
 FT CHAIN 1 550 HA2.
 SQ SEQUENCE 550 AA; 61802 MW; 1144331BC5A1F6A CRC64;

Query Match 92.3%; Score 132; DB 12; Length 550;

Best Local Similarity 95.7%; Pred. No. 1.6e-09;
Matches 22; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 GFGAAGIENGWEGMIDGWYG 24
|:|||||
Db 330 GFGAAGIENGWEGMIDGWYG 352

RESULT 15

Q82753 PRELIMINARY; PRT; 550 AA.
ID Q82753;
AC Q82753;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-OCT-2002 (TrEMBLrel. 22, Last annotation update)
DE Haemagglutinin (Fragment).
OS Influenza virus.
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC unclassified Orthomyxoviridae.
OX NCBI_TaxID=11309;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A/Philippines/2/82/BS/ML10;
RX MEDLINE=97300854; PubMed=9155874;
RA Hartley C.A., Reading P.C., Ward A.C., Anders E.M.;
RT "Changes in the haemagglutinin molecule of influenza type A (H3N2)
RL virus associated with increased virulence for mice.";
Arch. Virol. 142:75-88 (1997).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=A/Philippines/2/82/BS/ML10;
RX MEDLINE=97456249; PubMed=9311563;
RA Ward A.C.;
RT "Virulence of influenza A virus for mouse lung.";
RL Virus Genes 14:187-194 (1997).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
(HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR HMBP; U08905; AAC79579.1; -.
DR HSBP; P03437; 2VIU.
DR InterPro; IPR01364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
FT NON TER 1
FT CHAIN 1 328 HAEMAGGLUTININ HA1.
FT CHAIN 330 550 HAEMAGGLUTININ HA2.
SQ SEQUENCE 550 AA; 61745 MW; 692A49DB678AC4BC CRC64;

Query Match 92.3%; Score 132; DB 12; Length 550;
Best Local Similarity 95.7%; Pred. No. 1.6e-09;

Matches 22; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 GFGAAGIENGWEGMIDGWYG 24
|:|||||
Db 330 GFGAAGIENGWEGMIDGWYG 352

Search completed: January 30, 2004, 00:24:41
Job time : 26.7887 secs

A:Map position: 3
 A:Introns: 312/3; 359/3; 444/3
 C:Superfamily: Arabidopsis thaliana hypothetical protein F17J16.30

Query Match 48.8%; Score 102; DB 2; Length 517;
 Best Local Similarity 80.8%; Pred. No. 0.17;
 Matches 21; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

OY 20 EAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 45
 DB 440 ERVGGKKKKKKKKKKKKKKKKKKKKKK 465

RESULT 3

T39683
 znotin-like protein - fission yeast (Schizosaccharomyces pombe)

C:Species: Schizosaccharomyces pombe
 C:Date: 20-Oct-2000 #sequence_revision 20-Oct-2000 #text_change 20-Oct-2000
 C:Accession: T39683; T40195
 R:Oliver, K.; Harris, D.; Wood, V.; Rajandream, M.A.; Barrell, B.G.
 submitted to the EMBL Data Library, March 1998
 A:Reference number: 221869
 A:Accession: T39683
 A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA
 A:Residues: 1-124 <OLI>

A:Cross-references: EMBL:AL049489; PIDN:CA839796.1; GSPDB:GN00067; SPDB:SPBC1778.01c
 A:Experimental source: strain 972h-; cosmid c1778

R:Wood, V.; Rajandream, M.A.; Barrell, B.G.; Lauber, J.; Hiltbert, H.; Duesterhoeft, A.
 submitted to the EMBL Data Library, February 1998
 A:Reference number: 221910
 A:Accession: T40195

A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA

A:Residues: 89-442 <WOO>
 A:Cross-references: EMBL:Z97992; PIDN:CA810796.1; GSPDB:GN00067; SPDB:SPBC30D10.01
 A:Experimental source: strain 972h-; cosmid c30D10

C:Genetics:
 A:Gene: SPDB:SPBC1778.01c; SPDB:SPBC30D10.01
 A:Map position: 2

Query Match 48.6%; Score 101.5; DB 2; Length 442;
 Best Local Similarity 56.0%; Pred. No. 0.17;
 Matches 28; Conservative 3; Mismatches 12; Indels 7; Gaps 1;

OY 2 EAAAAA-----EAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 44
 DB 309 EAAAAAQQKKEEERRRAEAAAKASAAAANKKAKKQKRDKTVK 358

RESULT 4

152523
 nucleoporin p62 homolog - rat (fragment)

C:Species: Rattus norvegicus (Norway rat)
 C:Date: 27-Feb-1997 #sequence_revision 27-Feb-1997 #text_change 05-Nov-1999
 C:Accession: I52523
 R:Wang, Z.Q.; Akmal, K.M.; Kim, K.H.
 Biol. Reprod. 51, 1022-1030, 1994

A:Title: An unusual nucleoporin-related messenger ribonucleic acid is present in the ger
 A:Reference number: I52523; MUID:95151924; PMID:7849178
 A:Accession: I52523

A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: mRNA

A:Residues: 1-215 <RES>
 A:Cross-references: GB:S75997; NID:9913245; PIDN:AAB3384.1; PID:9913246
 A:Experimental source: testis

Query Match 47.8%; Score 100; DB 2; Length 215;
 Best Local Similarity 100.0%; Pred. No. 0.13;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 KKKKKKKKKKKKKKKKKKKKKKKKKKKKK 45
 |||||

DB 35 KKKKKKKKKKKKKKKKKKKKKKKKKKKKK 54

RESULT 5

T46395
 hypothetical protein DKFZp434i1120.1 - human (fragment)

C:Species: Homo sapiens (man)
 C:Date: 04-Feb-2000 #sequence_revision 04-Feb-2000 #text_change 04-Feb-2000
 C:Accession: T46395
 R:Ottenwajlder, B.; Obermayer, B.; Mewes, H.W.; Gassenhuber, J.; Wiemann, S.
 submitted to the Protein Sequence Database, January 2000
 A:Reference number: 223031
 A:Accession: T46395

A:Status: preliminary
 A:Molecule type: mRNA

A:Residues: 1-380 <AAA>
 A:Cross-references: EMBL:AL137556
 A:Experimental source: adult testis; clone DKFZp434i1120
 C:Genetics:

A:Note: DKFZp434i1120.1

Query Match 47.8%; Score 100; DB 2; Length 380;
 Best Local Similarity 100.0%; Pred. No. 0.2;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 KKKKKKKKKKKKKKKKKKKKKKKKKKKKK 45
 DB 355 KKKKKKKKKKKKKKKKKKKKKKKKKKKKK 374

RESULT 6

S09388
 histone H1 - sea urchin (Parechinus angulosus)

C:Species: Parechinus angulosus (angulate urchin)
 C:Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 24-Jul-1997
 C:Accession: S09388
 R:Hill, C.S.; Martin, S.R.; Thomas, J.O.
 EMBO J. 8, 2591-2599, 1989

A:Title: A stable alpha-helical element in the carboxy-terminal domain of free and chromo
 A:Reference number: S09388; MUID:90060019; PMID:2583125

A:Accession: S09388
 A:Status: preliminary

A:Molecule type: protein
 A:Residues: 1-206 <HIL>
 C:Superfamily: histone H1
 C:Keywords: chromosomal protein

Query Match 46.9%; Score 98; DB 2; Length 206;
 Best Local Similarity 55.8%; Pred. No. 0.18;
 Matches 24; Conservative 4; Mismatches 15; Indels 0; Gaps 0;

OY 2 EAAAAAEEAAAAEAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 44
 DB 140 KAAAKRKAAALAKKRAAAAKRAATKAKKPKKKTAKKAKK 182

RESULT 7

S25194
 znotin - yeast (Saccharomyces cerevisiae)

N:Alternate names: probable 2-DNA-binding protein; protein G9554; protein YGR285C
 C:Species: Saccharomyces cerevisiae
 C:Date: 28-May-1993 #sequence_revision 28-May-1993 #text_change 21-Jul-2000
 C:Accession: S25194; S64620; S19066
 R:Zheng, S.; Lockshin, C.; Herbert, A.; Winter, E.; Rich, A.
 EMBO J. 11, 3787-3796, 1992

A:Title: Znotin, a putative 2-DNA binding protein in Saccharomyces cerevisiae.
 A:Reference number: S25194; MUID:93010971; PMID:1396572

A:Accession: S25194
 A:Molecule type: DNA

A:Residues: 1-433 <ZNA>
 A:Cross-references: EMBL:X63612; NID:94836; PIDN:CA45156.1; PID:94837
 A:Note: part of this sequence, including the amino end of the mature protein, was confirm
 R:Voet, M.; Voelckaert, G.


```

C:Accession: F71619
R:Gardner, M.J.; Tetteelin, H.; Carucci, D.J.; Cummings, L.M.; Aravind, L.; Koonin, E.V.;
I:Perla, M.; Salzberg, S.; Zhou, L.; Sutton, G.G.; Clayton, R.; White, O.; Smith, H.O.
S:Science 282, 1126-1132, 1998
A:Title: Chromosome 2 sequence of the human malaria parasite Plasmodium falciparum.
A:Reference number: A71600; MUID:99021743; PMID:9804551
C:Accession: F71619
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-483 <GAP>
A:Cross-references: GB:AE001382; GB:AE001362; MID:93845130; PIDN:AC71836.1; PID:9384513
A:Experimental source: clone 3D7
C:Genetics:
A:Gene: PFB0235W

Query Match      43.1%   Score 90;   DB 2;   Length 483;
Best Local Similarity 85.0%;   Pred. No. 1.4;
Matches 17;   Conservative 3;   Mismatches 0;   Indels 0;   Gaps 0;

Cy 26 KKKKKKKKKKKKKKKKKKKKK 45
    |||||::|::|::|::|::|
Db 449 KKKKKKKKKKKKKKKKKKK 468

RESULT 13
T50609
hypothetical protein DKFZp761B2423.1 - human (fragment)
C:Species: Homo sapiens (man)
C:Date: 21-Jul-2000 #sequence_revision 21-Jul-2000 #text_change 21-Jul-2000
C:Accession: T50609
R:Blocker, H.; Boechat, M.; Brandt, P.; Mewes, H.W.; Weil, B.; Wiemann, S.
submitted to the Protein Sequence Database, June 2000
A:Reference number: Z25143
A:Accession: T50609
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-529 <AAA>
A:Cross-references: EMBL:AL359564
A:Experimental source: adult amygdala, clone DKFZp761B2423
C:Genetics:
A:Note: DKFZp761B2423.1

Query Match      43.1%   Score 90;   DB 2;   Length 529;
Best Local Similarity 45.0%;   Pred. No. 1.5;   11;   Indels 0;   Gaps 0;
Matches 18;   Conservative 11;   Mismatches 0;

Cy 5  AAATAAAATAAAATAAAKAKKKKKKKKKKKKKKKKK 44
    :::::|::|::|::|::|::|
Db 444 GSGGTRSHSSASASASODSKKKKKKKKKKKKKKKKK 483

RESULT 14
T23778
histone H1.1 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 04-Mar-2000
C:Accession: T23778; T42231; S09130; S01817
R:Percy, C.
submitted to the EMBL Data Library, August 1996
A:Reference number: Z19798
A:Accession: T23778
A:Status: translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-208 <WII>
A:Cross-references: GB:Z79603; PIDN:CAB01892.1; GSPDB:GN00028; CESP:ML63.3
A:Experimental source: clone M163
R:Jedrussik, M.; Schulze, E.
submitted to the EMBL Data Library, August 1997
A:Description: The histone H1 complement of Caenorhabditis elegans.
A:Reference number: Z22091
A:Accession: T42231
A:Status: translated from GB/EMBL/DBJ
A:Molecule type: mRNA

```

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A:Residues: 1-208 <JED>
A:Cross-references: EMBL:AF017810; PIDN:AAB70665.1
R:Sanicola, M.; Ward, S.; Childs, G.; Emmons, S.W.
U: Mol. Biol. 212, 259-268, 1990
A>Title: Identification of a Caenorhabditis elegans histone H1 gene family. Characterization
A:Reference number: 809130; PMID:90204554; PMID:1969492
A:Accession: S09130
A>Status: not compared with conceptual translation
A:Molecule type: DNA
A:Residues: 1-43,'T','45-83','H',85-100,'R',102-208 <SNA>
A:Cross-references: GB:X53277; NID:g10885; PIDN:CA37372.1; PID:g10886
R:Vanfleteren, J.R.; van Bun, S.M.; van Beeumen, J.J.
Biochem. J. 255, 647-652, 1988
A>Title: The primary structure of the major isoform (H1.1) of histone H1 from the nematode
A:Reference number: S01817; PMID:89076229; PMID:3202838
A:Accession: S01817
A:Molecule type: protein
A:Residues: 2-43,'T','45-100,'K',102-208 <VAN>
C:Genetics:
A:Gene: CESP:M163.3; his-24
A:Map position: X
A:Introns: 79/3
C/Superfamily: histone H1
C/Keywords: blocked amino end; chromosomal protein; DNA binding; nucleosome; nucleus
P:2-208/Product: histone H1.1 #status predicted <MKT>
F:2/Modified site: blocked amino end (Ser) (in mature form) (probably acetylated) #status
Query Match 42.6%; Score 89; DB 2; Length 208;
Best Local Similarity 53.5%; Pred. No. 0.92;
Matches 23; Conservative 4; Mismatches 16; Indels 0; Gaps 0;

```

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RESULT 15
519113
cgr-4 protein - Chlamydomonas reinhardtii (fragment)
C:Species: Chlamydomonas reinhardtii
C:Date: 13-Jan-1995 #sequence_revision 13-Jan-1995 #ext_change 21-Jul-2000
C:Accession: S19113; S14466
R:Wakarchuk, W.W.; Mueller, F.W.; Beck, C.F.
Plant Mol. Biol. 18, 143-146, 1992
A:Title: Two GC-rich DNA elements of Chlamydomonas reinhardtii with complex arrangements
A:Reference number: S19113; MOID:92119224; PMID:1731966
A:Accession: S19113
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-265 <MAX>
A:Cross-references: EMBL:X17208; NID:g18136; PIDN:CA45080.1; PID:g18137
C:Genetics:
A:Gene: cgr-4

Query Match          42.6%; Score 89; DB 2; Length 265;
Best local similarity 68.8%; Pred. No. 1.1;
Matches 22; Conservative 3; Mismatches 7; Indels 0; Gaps 0;

QY      3 AAAAAEAAAAAEAAAAAEAAAAAAXXXXXXXXXX 34
      ||| ||| ||| ||| ||| ||| ||| ||| |||
DB      176 AAAEAPAAAAAAEAAAAAAKAKAAAAAEAKKADK 207

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DB 146 KAAAKKAAALAKKAAAKKAAAKKAAAKKAAAKKAAAKKAAAKK 187

RESULT 2

Y011 SCHPO STANDARD; PRT; 442 AA.
AC 09y718: 014347;
ID 15-SEP-2003 (Rel. 42, Created)
DT 15-SEP-2003 (Rel. 42, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Hypothetical J-domain protein C1778.01c in chromosome II.
GN SPBC1778.01C OR SPBC30D10.01.
OS Schizosaccharomyces pombe (Fission yeast).
OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OC Schizosaccharomycetes.
OX NCBI_TaxID=4896;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=972;
RX MEDLINE=21849401; PubMed=11859360;
RA Wood V., Gilliam R., Rajandream M.A., Lyne M., Lyne R., Stewart A.,
RA Sgourou J., Peat N., Hayles J., Baker S., Basham D., Bowman S.,
RA Brooks K., Brown D., Brown S., Chillingworth T., Churcher C.M.,
RA Collins M., Connor R., Cronin A., Davis P., Fellwell T., Fraser A.,
RA Gentles S., Goble A., Hamlin N., Harris D., Hidalgo J., Hodgson G.,
RA Holroyd S., Hornby T., Howarth S., Huckle E.J., Hunt S., Jagsle K.,
RA James K., Jones L., Jones M., Leach S., McDonald S., McLean J.,
RA Mooney P., Moule S., Mungall K., Murphy L., Niblett D., Odell C.,
RA Oliver K., O'Neill S., Pearson D., Quail M.A., Rabinowitch E.,
RA Rutherford K., Rutter S., Saunders D., Seeger K., Sharp S.,
RA Skelton J., Simmonds M., Squares R., Squares S., Stevens K.,
RA Taylor K., Taylor R.G., Tivey A., Walsh S.V., Warren T., Whitehead S.,
RA Woodward J., Volkart G., Aert R., Robben J., Grynolprez B.,
RA Wellens J., Vanterreels B., Rieger M., Schaefer M., Mueller-Auer S.,
RA Gabel C., Fuchs M., Fritz C., Holzer E., Meest D., Hilbert H.,
RA Borzym K., Langer I., Beck A., Lehnach H., Reinhardt R., Pohl T.M.,
RA Egger P., Zimmermann W., Medler H., Wandut R., Purnelle B.,
RA Goffeau A., Cadieu E., Dreano S., Gloux S., Lelaure V., Mottier S.,
RA Gilbert F., Aves S.J., Xiang Z., Hunt C., Moore K., Huret S.M.,
RA Lucas M., Rochet M., Galliardin C., Tallada V.A., Garzon A., Thode G.,
RA Daga R.R., Cruzado L., Jimenez J., Sanchez M., del Rey F., Benito J.,
RA Dominguez A., Revuelta J.L., Moreno S., Armstrong J., Forsburg S.L.,
RA Cernuti L., Lowe T., McCombie W.R., Paulsen I., Potashkin J.,
RA Shpakovski G.V., Useery D., Barrett B.G., Nurse P.;
RT "The genome sequence of Schizosaccharomyces pombe.";
RL Nature 415:871-880(2002).
CC -1- SIMILARITY: Contains 1 J domain.
CC -----
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CC -----
CC EMBL; AL049489; CAB39796.1; -
DR EMBL; 297992; CAB10796.1; -
DR PIR; T39683; T39683.
DR HSSP; P25685; 1HDJ.
DR GeneDB; SPombe; SPBC1778.01c; -
DR InterPro; IPR001623; DnaJ_N.
DR Pfam; PF00226; DnaJ_1.
DR SMART; SM00271; DnaJ_1.
DR PROSITE; PS00636; DnaJ_1; 1.
DR PROSITE; PS00636; DnaJ_2; 1.
DR PROSITE; PS00636; DnaJ_2; 1.
KM Hypothetical protein; Chapone.
FT DOMAIN 97 167 J-DOMAIN.
FT DOMAIN 294 342 ALA/LYS-RICH.
SQ SEQUENCE 442 AA; 50209 MW; F4EC924871B7118B CRC64;

Query Match 48.6%; Score 101.5; DB 1; Length 442;
Best Local Similarity 56.0%; Pred. No. 0.042;
Matches 28; Conservative 3; Mismatches 12; Indels 7; Gaps 1;

RESULT 3

Y011 YEAST STANDARD; PRT; 433 AA.
AC P32527;
ID 01-OCT-1993 (Rel. 27, Created)
DT 01-OCT-1993 (Rel. 27, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE ZuoLin.
GN ZUO1 OR YGR285C.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A., AND SEQUENCE OF 1-10.
RC STRAIN=20B-12;
RX MEDLINE=93010971; PubMed=1396572;
RA Zhang S., Lockshin C., Herbert A., Winter E., Rich A.;
RA "ZuoLin, a putative Z-DNA binding protein in Saccharomyces
RT cerevisiae.";
RL EMBL J. 11:3787-3796(1992).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=8288C / FY1679;
RX MEDLINE=97245295; PubMed=9090054;
RA Volckaert G., Voet M., Robben J.;
RT "Sequence analysis of a near-subtelomeric 35.4 kb DNA segment on the
RT right arm of chromosome VII from Saccharomyces cerevisiae carrying
RT the MLI1 locus reveals 15 complete open reading frames, including
RT ZUO1, BGL2 and BHO2 genes and an ABC transporter gene.";
RL Yeast 13:251-259(1997).
CC -1- FUNCTION: Z-DNA BINDING PROTEIN. COULD BE INVOLVED IN
CC CHROMOSOME ORGANIZATION.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- SIMILARITY: Contains 1 J domain.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; X63612; CAA45156.1; -
DR EMBL; 273070; CAA97317.1; -
DR PIR; S25194; S25194.
DR HSSP; P08622; 1BQZ.
DR SGD; S0003517; ZUO1.
DR GO; GO:0005829; C:cytosol; NAS.
DR GO; GO:0005840; C:ribosome; NAS.
DR GO; GO:0003754; F:chaperone activity; NAS.
DR GO; GO:0006457; P:protein folding; NAS.
DR InterPro; IPR001623; DnaJ_N.
DR Pfam; PF00226; DnaJ_1.
DR SMART; SM00271; DnaJ_1.
DR PROSITE; PS00636; DnaJ_1; 1.
DR PROSITE; PS00636; DnaJ_2; 1.
KM Chaperone; DNA-binding; Nuclear protein.
FT DOMAIN 98 170 J-DOMAIN.
FT DOMAIN 306 357 ALA/LYS-RICH.
SQ SEQUENCE 433 AA; 49019 MW; 0AA76B1CD3C7DAB CRC64;
Query Match 45.9%; Score 96; DB 1; Length 433;

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Best Local Similarity 52.3%; Pred. No. 0.11; Indels 0; Gaps 0
Matches 23; Conservative 6; Mismatches 15; Indels 0; Gaps 0

QY      2 EAAAAAAAAAAAAAAAAAKKKKKKKKKKKKKK 45
       |||::|||::|||::|||::|||::|||::
Db       313 EAKAAEAEAKAASEAKANASAKADKKKAEEAKKKGR 356

RESULT 4
FA9A_HUMAN          STANDARD; PRT; 332 AA.
ID   FA9A_HUMAN     OGIUTL;
AC   OGIUTL;        DT 15-SEP-2003 (Rel. 42, Created)
DT 15-SEP-2003 (Rel. 42, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE   Protein FAM9A.
GN   FAM9A.
OS   Homo sapiens (Human).
OC   Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC   Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
OX   NCBI_TaxID=9606;
RN   [1]
RP   SEQUENCE FROM N.A., TISSUE SPECIFICITY, AND SUBCELLULAR LOCATION.
RX   MEDLINE=22202142; PubMed=12213195;
RA   Martinez-Garay I., Jablonka S., Sutajova M., Steuenkel P., Gal A.,
RA   Kutscie K.;
RT   "A new gene family (FAM9) of low-copy repeats in Xp22.3 expressed
RT   exclusively in testis: implications for recombinations in this
RT   region."
RL   Genomics 80:259-267(2002).
CC   -1- SUBCELLULAR LOCATION: Nuclear; nucleolar.
CC   -1- TISSUE SPECIFICITY: Expressed exclusively in testis.
CC   -1- SIMILARITY: Belongs to the FAM9 family.
-----
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CC   or send an email to license@sib-sib.ch).
-----
DR   EMBL; AF949433; AACD7162.1; -.
DR   Genew; HGNC:18403; FAM9A.
KW   Nuclear Protein.
FT   DOMAIN         180..275    GLU-RICH.
FT   FT             194..214    POLY-ALA.
FT   DOMAIN         252..258    POLY-GLY.
SQ   SEQUENCE       332 AA; 37339 MW; 92F22BC36038229C CRC64;

Query Match           44.5%; Score 93; DB 1; Length 332;
Best Local Similarity 44.2%; Pred. No. 0.16;
Matches 19; Conservative 17; Mismatches 7; Indels 0; Gaps 0

QY      3 AAAAAAAAAAAAAAAAAAKKKKKKKKKKKKKK 45
       |||::|||::|||::|||::|||::|||::
Db       197 AAAAAEAFAAAAALVIEDEBEEXKEEBEKEEBE 239

RESULT 5
H2B4_CHLRE          STANDARD; PRT; 153 AA.
ID   H2B4_CHLRE     PS4347;
AC   PS4347;        DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE   Histone H2B-IV.
OS   Chlamydomonas reinhardtii.
OC   Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
OC   Chlamydomonadaceae; Chlamydomonas.
OX   NCBI_TaxID=3055;
RN   [1]
RP   SEQUENCE FROM N.A.
```

```

RX MEDLINE=96120862; PubMed=8590479;
RA Fabry S., Mueller K., Lindauer A., Park P.B., Cornelius T.,
RA Schmitt R.;
RT "The organization structure and regulatory elements of Chlamydomonas
RT histone gene reveal features linking plant and animal genes.";
RL Curr. Genet. 28:333-345(1995).
CC
CC -1- SUBUNIT: The nucleosome is an octamer containing two molecules
CC each of H2A, H2B, H3 and H4. The octamer wraps approximately 146
CC bp of DNA.
CC
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC
CC -1- SIMILARITY: Belongs to the histone H2B family.
CC
CC -----
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CC -----
CC EMBL; U16726; AAA98454.1; -.
CC PIR; S59591; S59591.
CC InterPro: IPR004822; Histone_core.
CC InterPro: IPR00558; Histone_H2B.
CC Pfam; PF00125; histone; 1.
CC PRINTS; PR00621; HISTONEH2B.
CC PRODOM; PD000497; Histone_H2B; 1.
CC SMART; SM00427; H2B; 1.
CC POSITE; PS00357; HISTONE_H2B; 1.
KW Chromosomal protein; Nucleosome core; Nuclear protein; DNA-binding;
KW Multigene family.
KW SEQUENCE 153 AA; 16587 MW; B09CE1D804E1F485 CRC64;
SQ
Query March 43.1%; Score 90; DB 1; Length 153;
Best Local Similarity 51.0%; Pred. No. 0.14;
Matches 25; Conservative 2; Mismatches 16; Indels 6; Gaps 1;
QY 3 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAKKKKKKKKKK-----KKKKKKKK 45
Db 12 AEAQAEPAPAKAEKPKAKAKKAKKPKSPSKAKAEKPGDKDKKKKK 60
RESULT 6
H1 CAEEL STANDARD; PRT; 207 AA.
ID ID_H1 CAEEL
AC P10771;
DT 01-JUN-1989 (Rel. 11, Created)
DT 01-AUG-1991 (Rel. 19, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE Histone H1.1.
GN H1S-24.
OS Caenorhabditis elegans.
OC Eukaryotes; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
OC Rhabditiidae; Peloiderinae; Caenorhabditis.
OX NCB1 TaxID=6239;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90204554; PubMed=1969492;
RA Sanicola M., Ward S., Childs G., Emmons S.W.;
RT "Identification of a Caenorhabditis elegans histone H1 gene family.
RT Characterization of a family member containing an intron and encoding
RT a poly(A)+ mRNA.";
RL J. Mol. Biol. 212:259-268(1990).
RN [2]
RP SEQUENCE.
RC STRAIN=Bristol N2;
RX MEDLINE=89076229; PubMed=3202839;
RA Vanfleteren J.R., Van Bun S.M., Van Beumen J.J.;
RT "The primary structure of the major isoform (H1.1) of histone H1 from
RT the nematode Caenorhabditis elegans.";
RL Biochem. J. 255:647-652(1988).
CC -1- FUNCTION: HISTONES H1 ARE NECESSARY FOR THE CONDENSATION OF
CC NUCLEOSOME CHAINS INTO HIGHER ORDER STRUCTURES.

```

CC -1- SUBCELLULAR LOCATION: Nuclear.
 CC -1- MISCELLANEOUS: THERE ARE 2 FORMS OF H1 IN THIS NEMATODE: H1.1 IS
 CC THE MAJOR FORM.
 CC -1- SIMILARITY: BELONGS TO THE HISTONE H1/H5 FAMILY.
 CC -----
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 CC -----
 DR EMBL: X53277; CAA37372.1; -.
 DR HSSP: P02259; H1ST.
 DR InterPro: IPR005818; Histone_H1/H5.
 DR InterPro: IPR005819; Histone_H5.
 DR InterPro: IPR003216; Linkerhist_N.
 DR Pfam: PF00538; linker histone; 1.
 DR PRINTS: PR00624; HISTONEH5.
 DR ProDom: PD000373; Linkerhist_N; 1.
 DR SMART: SM00525; H15; 1.
 DR Chromosomal protein; Nuclear protein; DNA-binding; Multigene family;
 KM Acetylation.
 FT INIT MET 0 0
 FT MOD RES 1 1 ACETYLATION (PROBABLE).
 FT DOMAIN 36 112 GLOBULAR.
 FT CONFLICT 83 83 H -> L (IN REF. 2).
 FT CONFLICT 100 100 R -> K (IN REF. 2).
 SQ SEQUENCE 207 AA; 21314 MW; 7802EA9AA4F36F6F CRC64;

Query Match 42.6%; Score 89; DB 1; Length 207;
 Best Local Similarity 53.5%; Pred. No. 0.23; 16; Indels 0; Gaps 0;
 Matches 23; Conservative 4; Mismatches 16; Indels 0; Gaps 0;

QY 2 EAA 44
 DB 112 EKAATAKKPAKKPAKKPAKKPAKKPAKKPAKKPAKKPAKKPAKKPAKK 154

RESULT 7
 TOLA_PSEAE STANDARD; PRT; 347 AA.
 AC P50600;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE TOLA protein.
 GN TOLA OR PA0971.
 OS Pseudomonas aeruginosa.
 CC Bacteria; Proteobacteria; Gammaproteobacteria; Pseudomonadales;
 CC Pseudomonadaceae; Pseudomonas.
 OX NCBI_TaxID=287;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=PAO;
 RX MEDLINE=97113525; PubMed=8955385;
 RA Dennis J.J., Lafontaine E.R., Sokol P.A.;
 RT "Identification and characterization of the *tolJ* genes of
 RT *Pseudomonas aeruginosa*.";
 RL J. Bacteriol. 178:7059-7068(1996).
 RN [2]
 RP REVISIONS TO N-TERMINUS.
 RA Duan X., Sokol P.A.;
 RL Submitted (Aug-1999) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC 15692 / PAO1;
 RX MEDLINE=20437337; PubMed=10984043;
 RA Stever C.K., Pham X.-Q.T., Erwin A.L., Mizoguchi S.D., Warren P.,
 RA Hickey M.J., Brinman F.S.L., Hufnagle W.O., Kowalik D.J., Lagrou M.,
 RA Garner R.L., Goltzy L., Tolentino E., Westbrock-Wadman S., Yuan Y.,
 RA Brody L.L., Coulter S.N., Folger K.R., Kas A., Lartig K., Lim R.M.,

RA Smith K.A., Spencer D.H., Wong G.K.-S., Wu Z., Paulsen I.T.,
 RA Reizer J., Sailer M.H., Hancock R.E.W., Lory S., Olson M.V.;
 RT "Complete genome sequence of *Pseudomonas aeruginosa* PAO1, an
 RT opportunistic pathogen.";
 RL Nature 406:599-964(2000).
 CC -1- FUNCTION: INVOLVED IN THE TONB-INDEPENDENT UPTAKE OF PROTEINS
 CC (BY SIMILARITY).
 CC -1- SUBCELLULAR LOCATION: Type II membrane protein. Inner membrane
 CC (potential).
 CC -----
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 CC -----
 DR EMBL: U39558; AAC4460.2; -.
 DR EMBL: AB004530; AAG04360.1; -.
 DR PIR: B83525; B83525.
 DR InterPro: IPR006260; TonB_C.
 DR TIGRPFAMs: TIGR01352; TonB_Cterm; 1.
 KM Transport; Protein transport; Transmembrane; Repeat; Inner membrane;
 KM Complete proteome.
 FT DOMAIN 1 16 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 17 37 POTENTIAL.
 FT DOMAIN 38 347 PERIPLASMIC (POTENTIAL).
 FT POLY-ALA 209 216 POLY-ALA.
 SQ SEQUENCE 347 AA; 37935 MW; EEDDA4B0A095945 CRC64;

Query Match 41.1%; Score 86; DB 1; Length 347;
 Best Local Similarity 51.2%; Pred. No. 0.61; 16; Indels 0; Gaps 0;
 Matches 22; Conservative 5; Mismatches 16; Indels 0; Gaps 0;

QY 2 EAA 44
 DB 124 EAQAAAEAKKDAEAKKAFAKAEKQADYAKRAEDAEAKK 166

RESULT 8
 RL12_METUA
 ID RL12_METUA STANDARD; PRT; 102 AA.
 AC P54048;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 16-OCT-1996 (Rel. 34, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE 50S ribosomal protein L12P.
 GN RPL12P OR M0508.
 OS Methanococcus jannaschii.
 CC Archaea; Euryarchaeota; Methanococci; Methanococcales;
 CC Methanocaldococcaceae; Methanocaldococcus.
 OX NCBI_TaxID=2190;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=JAL-1 / DSM 2661 / ATCC 43067;
 RX MEDLINE=96337999; PubMed=8688087;
 RA Bult C.J., White O., Olsen G.J., Zhou L., Fleischmann R.D.,
 RA Sutton G.G., Blake J.A., Fitzgerald L.M., Clayton R.A., Gocayne J.D.,
 RA Kerlavage A.R., Dougherty B.A., Tomb J.-F., Adams M.D., Reich C.I.,
 RA Overbeek R., Kirkness E.F., Weinstock K.G., Merrick J.M., Glodek A.,
 RA Scott J.L., Geophagen N.S.M., Weidman J.F., Fuhrmann J.L., Nguyen D.,
 RA Uitterback T.R., Kelley J.M., Peterson J.D., Sadow P.W., Hanna M.C.,
 RA Cotton W.D., Roberts K.M., Huret M.A., Kaine B.P., Borodovsky M.,
 RA Kleink H.-P., Frazer C.W., Smith H.O., Weese C.R., Venter J.C.;
 RT "Complete genome sequence of the methanogenic archaeon, *Methanococcus*
 RT *jannaschii*.";
 RL Science 273:1058-1073(1996).
 CC -1- FUNCTION: SEEMS TO BE THE BINDING SITE FOR SEVERAL OF THE FACTORS
 CC INVOLVED IN PROTEIN SYNTHESIS AND APPEARS TO BE ESSENTIAL FOR
 CC ACCURATE TRANSLATION (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE L12P FAMILY OF RIBOSOMAL PROTEINS.
 CC -----

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DR EMBL; U67500; AAB98498.1; -
DR PIR; D64363; D64363.
DR TIGR; M0508; -
DR InterPro; IPR001813; 60s_ribosomal.
DR Pfam; PF00428; 60s_ribosomal; 1.
KW Ribosomal protein; Complete proteome.
SQ SEQUENCE 102 AA; 10363 MW; 35506FDE967C52C CRC64;

Query Match 40.7%; Score 85; DB 1; Length 102;
Best Local Similarity 52.6%; Pred. No. 0.26;
Matches 20; Conservative 8; Mismatches 10; Indels 0; Gaps 0;

Db 2 EAAAAAAAAAAAAAAAAAKKKKKKKKKKKK 39
52 EAIANAMPVAAAPAAAAAERKKEKKEKKEK 89

RESULT 9
ID H2B3_CHLRE STANDARD; PRT; 153 AA.
AC P54346;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Histone H2B-III.
OS Chlamydomonas reinhardtii.
OC Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
OC Chlamydomonadaceae; Chlamydomonas.
OX NCBI_TaxID=3055;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=96120862; PubMed=8590479;
RA Fabry S., Mueller K., Lindner A., Park P.B., Cornelius T.,
RA Schmitt R.;
RT "The organization structure and regulatory elements of Chlamydomonas
RT histone genes reveal features linking plant and animal genes";
RL Curr. Genet. 28:333-345(1995).
CC -1- SUBUNIT: The nucleosome is an octamer containing two molecules
CC each of H2A, H2B, H3 and H4. The octamer wraps approximately 146
CC BP of DNA.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- SIMILARITY: Belongs to the histone H2B family.
CC -----
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DR EMBL; U16725; AAA98450.1; -
DR PIR; S59587; S59587.
DR InterPro; IPR004822; Histone_core.
DR InterPro; IPR00558; Histone_H2B.
DR Pfam; PF00125; histone; 1.
DR PRINTS; PR00621; HISTONEH2B.
DR ProDom; PD000497; Histone_H2B; 1.
DR SMART; SM00427; H2B; 1.
DR PROSITE; PS00357; HISTONE_H2B; 1.
KW Chromosomal protein; Nucleosome core; Nuclear protein; DNA-binding;
KW Multigene family.
SQ SEQUENCE 153 AA; 16557 MW; 2092413E04E1F49C CRC64;

Query Match 40.7%; Score 85; DB 1; Length 153;

Best Local Similarity 49.0%; Pred. No. 0.37;
Matches 24; Conservative 2; Mismatches 17; Indels 6; Gaps 1;

Db 3 AAAAAAAAAAAAAAAAAAKKKKKKKKKK-----KKKKKKKK 45
12 AEAAGAAAPAAKAPKAKGKAKKAPKAAKAPKGGKKKKKK 60

RESULT 10
ID H12_VOLCA STANDARD; PRT; 240 AA.
AC Q08865;
DT 01-OCT-1994 (Rel. 30, Created)
DT 01-FEB-1995 (Rel. 31, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Histone H1-II.
OS H1-II.
OS Volvox carteri.
OC Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
OC Volvocaceae; Volvox.
OX NCBI_TaxID=3067;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=f. Nagariensis / HK10;
RX MEDLINE=93328125; PubMed=8335260;
RA Lindner A., Mueller K., Schmitt R.;
RT "Two histone H1-encoding genes of the green alga Volvox carteri with
RT features intermediate between plant and animal genes.";
RL Gene 129:59-68(1993).
CC -1- FUNCTION: HISTONES H1 ARE NECESSARY FOR THE CONDENSATION OF
CC NUCLEOSOME CHAINS INTO HIGHER ORDER STRUCTURES.
CC -1- SUBCELLULAR LOCATION: Nuclear.
CC -1- DEVELOPMENTAL STAGE: EXPRESSION IS RESTRICTED TO EMBRYOGENESIS.
CC -1- SIMILARITY: BELONGS TO THE HISTONE H1/H5 FAMILY.
CC -----
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DR EMBL; L07947; AAA34246.1; -
DR PIR; JN0748; JN0748.
DR InterPro; IPR005818; Histone_H1/H5.
DR InterPro; IPR005819; Histone_H5.
DR InterPro; IPR003216; Linkerhist_N.
DR Pfam; PF00538; linker histone; 1.
DR PRINTS; PR00624; HISTONEH5.
DR ProDom; PD000373; Linkerhist_N; 1.
DR SMART; SM00526; H15; 1.
KW Chromosomal protein; Nuclear protein; DNA-binding; Multigene family;
KW Repeat.
FT INIT MET 0 0 BY SIMILARITY.
FT DNA_BIND 21 98 POTENTIAL.
FT DNA_BIND 182 185 POTENTIAL.
FT DNA_BIND 202 205 POTENTIAL.
FT DNA_BIND 210 213 POTENTIAL.
FT DOMAIN 110 216 8 X 6 AA REPEATS OF P-K-K-A-[AK]-A.
FT REPEAT 110 115 1.
FT REPEAT 116 121 1.
FT REPEAT 122 127 2.
FT REPEAT 128 133 3.
FT REPEAT 133 138 4.
FT REPEAT 138 143 5.
FT REPEAT 143 148 6.
FT REPEAT 148 153 7.
FT REPEAT 153 158 8.
FT REPEAT 158 163 7.
FT REPEAT 163 168 6.
FT REPEAT 168 173 5.
FT REPEAT 173 178 4.
FT REPEAT 178 183 3.
FT REPEAT 183 188 2.
FT REPEAT 188 193 1.
FT REPEAT 193 198 0.
FT REPEAT 198 203 0.
FT REPEAT 203 208 0.
FT REPEAT 208 213 0.
FT REPEAT 213 216 0.
SQ SEQUENCE 240 AA; 25072 MW; 01AA389E08F421BD CRC64;

Query Match 40.7%; Score 85; DB 1; Length 240;
Best Local Similarity 50.0%; Pred. No. 0.54;
Matches 23; Conservative 7; Mismatches 14; Indels 2; Gaps 1;

```

QY      2 EAAAAAEAAAAAEAAAAAAXXXXXXXXXXXXXXXXXX--XXXXXXXX 45
          : ||| : ||| : ||| ||| ||| : ||| ||| : |||
Db      99 KAKAAAKPKAAPKCAAAAPKCAAAAPKCAKAPKKEGKKA VKPKSEKK 144

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ID	PHPA_PLACH	STANDARD;	PRT;	441 AA.
AC	002752;			
DT	01-JUL-1993 (Rel. 26, Created)			
DT	01-JUL-1993 (Rel. 26, Last sequence update)			
DT	01-JUN-1994 (Rel. 29, Last annotation update)			
DE	Acidic phosphoprotein precursor (50 kDa antigen).			
GN	PCEM1.			
OS	Plasmodium chabaudi.			
OC	Eukaryote; Alveolata; Apicomplexa; Haemosporida; Plasmodium.			
OX	NCBI_TaxID=35825;			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	STRAIN=IP-PCI;			
RX	MEDLINE=93116806; PubMed=1475002;			
RA	Deleensijder W., Prasomsitti P., Tungpradubkul S., Hendrix D.,			
RA	Hamers-Casterman C., Hamers R.;			
RT	"Structure of a Plasmodium chabaudi acidic phosphoprotein that is			
RT	associated with the host erythrocyte membrane.";			
RL	Mol. Biochem. Parasitol. 56:59-68(1992).			
CC	-1- FUNCTION: DURING INFECTION, THIS PHOSPHOPROTEIN PROBABLY MODULATES			
CC	THE STRUCTURE OF THE RED CELL MEMBRANE TO THE ADVANTAGE OF THE			
CC	PARASITE. ALTHOUGH ITS PRECISE FUNCTION IS NOT KNOWN, THE			
CC	-1- SUBCELLULAR LOCATION: PERIPHERAL MEMBRANE PROTEIN ON THE			
CC	CYTOPLASMIC FACE OF THE HOST ERYTHROCYTE MEMBRANE.			
CC	-1- MISCELLANEOUS: ASSOCIATED WITH THE HOST RED CELL MEMBRANE			
CC	THROUGHOUT THE ENTIRE ERYTHROCYTIC CYCLE.			
CC	-----			
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CC	-----			
DR	EMBL; M95769; AAA29732.1; -.			
DR	PIR; A48455; A48455.			
KW	Phosphorylation; Signal; Antigen; Membrane; Repeat; Erythrocyte.			
FT	CHAIN	1	15	OR 24 (POTENTIAL).
FT	DOWAIN	16	441	ACIDIC PHOSPHOPROTEIN.
FT	REPEAT	186	193	16 X 8 AA TANDEM REPEATS.
FT	REPEAT	194	201	1-1.
FT	REPEAT	202	209	1-2.
FT	REPEAT	210	217	1-3.
FT	REPEAT	218	225	1-4.
FT	REPEAT	226	233	1-5.
FT	REPEAT	234	241	1-6.
FT	REPEAT	242	249	1-7.
FT	REPEAT	250	257	1-8.
FT	REPEAT	258	265	1-9.
FT	REPEAT	266	273	1-10.
FT	REPEAT	274	281	1-11.
FT	REPEAT	282	289	1-12.
FT	REPEAT	290	297	1-13.
FT	REPEAT	298	305	1-14.
FT	REPEAT	306	313	1-15.
FT	DOWAIN	353	370	1-16.
FT	REPEAT	353	360	2 X 9 AA TANDEM REPEATS.
FT	REPEAT	361	368	2-1.
FT	DOWAIN	371	417	2-2.
FT	CARBOHYD	21	21	LYS-RICH (BASIC).
FT	CARBOHYD	112	112	N-LINKED (GLCNAC. . .) (POTENTIAL).
QO	SEQUENCE	441 AA;	49708 MW;	N-LINKED (GLCNAC. . .) (POTENTIAL).

Query Match	40.2%	Score 84;	DB 1;	Length 441;
Best Local Similarity	80.0%	Pred. No. 1.1;		
Matches 16;	Conservative 4;	Mismatches 0;	Indels 0;	Gaps 0;

```
QY      26  KKKKKKKKKKKKKKKKKKKKK 45
          |||||:|||||:|:||||:
Db     394  KKKKKKKKKKKKKKKKKKK 413
```

ID	BRD3 HUMAN	STANDARD;	PRT;	726 AA.
AC	Q15059; Q92645;			
DT	16-OCT-2001 (Rel. 40, Created)			
DT	16-OCT-2001 (Rel. 40, Last sequence update)			
DT	16-OCT-2001 (Rel. 40, Last annotation update)			
DE	Bromodomain-containing protein 3 (RING3-like protein).			
GN	BRD3 OR RING3L OR KIAA0043.			
OS	Homo sapiens (Human).			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
OC	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
OX	NCBI_TaxId=9606;			
XP	[1]			
RP	SEQUENCE FROM N.A.			
RC	TISSUE=Bone marrow;			
RX	MEDLINE=96051398; PubMed=7584044;			
RA	Nomura N., Nagase T., Miyajima N., Sazuka T., Tanaka A., Sato S.,			
RA	Seki N., Kawabayashi Y., Ishikawa K.-I., Tabata S.;			
RT	"Prediction of the coding" sequences of unidentified human genes. II.			
RT	The coding sequences of 40 new genes (KIAA0041-KIAA0080) deduced by			
RT	analysis of cDNA clones from human cell line KG-1."			
RL	DNA Res. 1:223-229(1994).			
RP	[2]			
RX	SEQUENCE OF 363-726 FROM N.A.			
RA	MEDLINE=98038990; PubMed=9373153;			
RA	Thorpe K.L., Gorman P., Thomas C., Sheer D., Trowsdale J., Beck S.;			
RT	"Chromosomal localization, gene structure and transcripion pattern of			
RT	the ORFX gene, a homologue of the MHC-linked RING3 gene."			
RL	Gene 200:177-183(1997).			
CC	-1- SUBCELLULAR LOCATION: Nuclear (potential).			
CC	-1- TISSUE SPECIFICITY: Ubiquitous.			
CC	-1- SIMILARITY: Contains 2 bromodomains.			
CC	-----			
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CC	entities requires a license agreement (See http://www.isb-sib.ch/announce/			
CC	or send an email to license@isb-sib.ch).			
CC	-----			
DR	EMBL; D26362; BAA05393.1; -.			
DR	EMBL; Z81330; CAB03630.1; -.			
DR	HSSP; Q92831; 1B91.			
DR	Genew; HGNC:1104; BRD3.			
DR	MTM; 601541; -.			
DR	GO; GO:0005634; C:nucleus; NAS.			
DR	InterPro; IPR001487; Bromodomain.			
DR	Pfam; PF00439; bromodomain; 2.			
DR	PRINTS; PRO0503; BROMODOMAIN.			
DR	SMART; SMD0297; BROMO. 2.			
DR	PROSITE; PS00633; BROMODOMAIN 1; 2.			
DR	PROSITE; PS50014; BROMODOMAIN 2; 2.			
KW	Bromodomain; Repeat; Nuclear protein.			
FT	DOMAIN 56 115 BROMODOMAIN 1.			
FT	DOMAIN 326 398 BROMODOMAIN 2.			
FT	DOMAIN 487 555 LYS-RICH.			
FT	DOMAIN 676 725 SER-RICH.			
FT	CONFLICT 465 466 EL -> DV (IN REF. 2).			
SO	SEQUENCE 726 AA; 79541 MW; 64F526FC3C1033AA CRC64;			
Query Match	40.2%;	Score 84;	DB 1;	Length 726;
Best Local Similarity	54.3%;	Pred. No. 1.7;		

Matches 19; Conservative 4; Mismatches 12; Indels 0; Gaps 0;

QY 11 AAAAAAAAAAAAAAAAAKKKKKKKKKKKKKKKKKKKK 45
DB 472 AVHEQALASQAPVKKKKKKKKKKKKKKKKKKKK 506

RESULT 13

ID_RL12_PVRAB STANDARD; PRT; 107 AA.

AC Q9UXS6;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE 50S ribosomal protein L12P
GN RPL12P OR PYRAB17820 OR PAB1168.
OS Pyrococcus abyssi.
OC Archaea; Euryarchaeota; Thermococci; Thermococcales; Thermococcaceae;
OC Pyrococcus.
OX NCBI_TaxID=29292;

[1] SEQUENCE FROM N.A.
RC STRAIN=GES / Orsay;
RX PubMed=12622808;

RA Cohen G.N., Barbe V., Flament D., Galperin M., Hellig R., Lecompte O.,
Voch O., Prieur D., Querellou J., Ripp R., Thierry J.-C.,
Van der Oost U., Weissenbach J., Zivanovic Y., Forterre P.;
RT "An integrated analysis of the genome of the hyperthermophilic
archaeon Pyrococcus abyssi.";
RL Mol. Microbiol. 47:1495-1512(2003).
CC -1- FUNCTION: SEEMS TO BE THE BINDING SITE FOR SEVERAL OF THE FACTORS
INVOLVED IN PROTEIN SYNTHESIS AND APPEARS TO BE ESSENTIAL FOR
ACCURATE TRANSLATION (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE L12P FAMILY OF RIBOSOMAL PROTEINS.

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or send an email to license@isb-sib.ch).

DR EMBL; AJ248288; CAB50687.1; -.
DR PIR; A75031; A75031.
DR InterPro; IPR001813; 60S_ribosomal.
DR Pfam; PF00428; 60S_ribosomal; 1.
KW Ribosomal protein; Complete proteome.
SQ SEQUENCE 107 AA; 11300 MW; CD423930BAD4ED CRC64;

Query Match 39.7%; Score 83; DB 1; Length 107;
Best Local Similarity 44.2%; Pred. No. 0.39;
Matches 19; Conservative 12; Mismatches 12; Indels 0; Gaps 0;

QY 2 EAAAAAAAAAAAAAAAAKKKKKKKKKKKKKKKKKKKK 44
DB 55 EGAAMPVAAPAAAPAPAEKKEKKEKKEKKEEVESE 97

RESULT 14

ID_YD33_YEAST STANDARD; PRT; 320 AA.

AC Q12117;
DT 30-MAY-2000 (Rel. 39, Created)
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hypothetical 36.2 kDa protein in RAD28-LYS14 intergenic region.
GN YDR033W OR YD9673.03
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
OX NCBI_TaxID=4932;

[1] SEQUENCE FROM N.A.
RC STRAIN=K12 / MG1655;
RX MEDLINE=97426617; PubMed=9278503;

RP SEQUENCE FROM N.A.
RA Arnold W., Becker A., Jaeger W., Kuester H., Nussbaumer B.;
RL Submitted (JUL-1996) to the EMBL/Genbank/DBJ databases.

RP SEQUENCE FROM N.A.
RC STRAIN=S288c / AB972;
RA Connor R., Churcher C.M., Barrell B.G., Rajandream M.A., Walsh S.V.;
RL Submitted (DEC-1995) to the EMBL/Genbank/DBJ databases.
CC -1- SUBCELLULAR LOCATION: Integral membrane protein (Probable).
CC -1- SIMILARITY: BELONGS TO THE ARCHAEAL OPSIN FAMILY. HSP30
SUBFAMILY.

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DR EMBL; Z74329; CAA98855.1; -.
DR EMBL; Z68196; CAA92370.1; -.
DR PIR; S61586; S61586.
DR SGI; S0002440; YDR033W.
DR GO; GO:0005886; C:plasma membrane; IDA.
DR InterPro; IPR001425; Bac_rhodopsin.
DR Pfam; PF01036; Bac_rhodopsin; 1.

KW Hypothetical protein; Transmembrane.
FT TRANSMEM 35 55 POTENTIAL.
FT TRANSMEM 63 83 POTENTIAL.
FT TRANSMEM 117 137 POTENTIAL.
FT TRANSMEM 142 162 POTENTIAL.
FT TRANSMEM 168 188 POTENTIAL.
FT TRANSMEM 205 225 POTENTIAL.
FT TRANSMEM 239 259 POTENTIAL.
FT DOMAIN 300 318 LYS-RICH.
SQ SEQUENCE 320 AA; 36190 MW; 4311F64B6AA209F CRC64;

Query Match 39.7%; Score 83; DB 1; Length 320;
Best Local Similarity 48.8%; Pred. No. 1;
Matches 21; Conservative 4; Mismatches 14; Indels 4; Gaps 1;

QY 7 AAAAAAAAAAAAAAAAAAKKKKKKKKKKKKKKKKKKK 45
DB 277 AEAAPAPVAPVAPPAATPNLSKDKKKSKSKSKSKSE 319

RESULT 15

ID_TOLA_ECOLI STANDARD; PRT; 421 AA.

AC P19934;
DT 01-FEB-1991 (Rel. 17, Created)
DT 01-FEB-1991 (Rel. 17, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE TOLA OR CIM OR EXCC OR LKY OR B0739.
GN TOLA OR CIM OR EXCC OR LKY OR B0739.
OS Escherichia coli.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Escherichia.

OX NCBI_TaxID=562;
[1] SEQUENCE FROM N.A.
RC STRAIN=JM105;
RX MEDLINE=90078104; PubMed=2687247;
RT Levensgood S.K., Webster R.E.;
RT "Nucleotide sequences of the tola and tolB genes and localization of
their products components of a multistep translocation system in
Escherichia coli.";
RT J. Bacteriol. 171:6600-6609(1989).
RN [2] SEQUENCE FROM N.A.
RC STRAIN=K12 / MG1655;
RX MEDLINE=97426617; PubMed=9278503;

RESULT 2
 Q8INH9 PRELIMINARY; PRT; 2347 AA.
 ID Q8INH9;
 AC Q8INH9;
 DT 01-MAR-2003 (TREMBLrel. 23, Created)
 DT 01-MAR-2003 (TREMBLrel. 23, Last sequence update)
 DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
 DE CG7518-PB.
 GN CG7518.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydriidae; Drosophilidae; Drosophila.
 OC NCBI_taxid=7227;
 (1)
 RP SEQUENCE FROM N.A.
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galie R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutcliffe G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.H., Blazer R.G., Champe M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Gabor L.D.,
 RA Abril J.F., Agbayani A., An H.J., Andrews-Pfankuch C., Baldwin D.,
 RA Bailew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Bortova D., Botchan M.R., Bouck J., Brokstein P., Brothier P.,
 RA Burris K.C., Busan D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
 RA Foster C., Gabrielian A.E., Gary N.S., Gelbart W.M., Glaeser K.,
 RA Glodex A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.J., Mei M.H., Ibegwan C.,
 RA Jatali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Laslo P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,
 RA Merulov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
 RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
 RA Nelson D.R., Nelson K.A., Nixon K., Nuskern D.R., Pacle J.M.,
 RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puti V., Reese M.G.,
 RA Reinert K., Remington K., Saunders R.D., Scheeler F., Shen H.,
 RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
 RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
 RA Svitskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
 RA Wang Z.Y., Waastman D.A., Weinstein G.M., Weissbach J.,
 RA Williams S.M., Woodgett W., Worley K.C., Wu D., Yang S., Yao Q.A., Ye J.,
 RA Yeh R.F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
 RA Zheng X.H., Zhong F.N., Zhou W., Zhou X., Zhu S., Zhu X., Smith H.O.,
 RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
 RT "The genome sequence of Drosophila melanogaster.";
 RL Science 287:2185-2195(2000).
 (12)
 RP SEQUENCE FROM N.A.
 RA Celniker S.E., Adams M.D., Krommiller B., Wan K.H., Holt R.A.,
 RA Evans C.A., Gocayne J.D., Amanatides P.G., Brandon R.C., Rogers Y.,
 RA Barton J., An H., Baldwin D., Barton J., Beeson K.Y., Busan D.A.,
 RA Carlson J.M., Center A., Champe M., Davenport L.B., Dietz S.M.,
 RA Dodson K., Dorett V., Doup L.E., Doyle C., Dresnek D., Fartan D.,
 RA Ferreira S., Frise E., Galie R.F., Gary N.S., George R.A.,
 RA Gonzalez M., Houck J., Hoskins R.A., Hostin D., Howland T.J.,
 RA Ibegwan C., Jatali M., Kruse D., Li P., Mattei B., Moshrefi A.,
 RA McIntosh T.C., Moy M., Murphy B., Nelson C.R., Nunoo J.,
 RA Pacle J., Paragay V., Park S., Patel S., Pfeiffer B.,
 RA Phouanavong S., Pittman G.S., Puti V., Richards S., Scheeler F.,
 RA Stapleton M., Strong R., Svitskas R., Tector C., Tyler D.,
 RA Williams S.M., Zaveri J.S., Smith H.O., Venter J.C., Rubin G.M.,
 RT "Sequencing of Drosophila melanogaster genome.";

RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 (3)
 RP SEQUENCE FROM N.A.
 RA Miera S., Crosby M.A., Matthews B.B., Bayraktaroglu L., Campbell K.,
 RA Hradecky P., Huang Y., Kaminker J.S., Prochnik S.B., Smith C.D.,
 RA Tupy J.L., Bergman C., Berman B., Carlson J.W., Celniker S.E.,
 RA Clamp M., Drysdale R., Emmert D., Frise E., de Grey A., Harris N.,
 RA Krommiller B., Marshall B., Millburn G., Richter J., Russo S.,
 RA Seale S.M.J., Smith E., Shu S., Smutnick F., Whitfield E.,
 RA Ashburner M., Gelbart W.M., Rubin G.M., Mungall C.J., Lewis S.E.;
 RT "Annotation of Drosophila melanogaster genome.";
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 (4)
 RP SEQUENCE FROM N.A.
 RA Adams M.D., Celniker S.E., Gibbs R.A., Rubin G.M., Venter J.C.;
 RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
 (5)
 RP SEQUENCE FROM N.A.
 RA FLYBASE;
 RP Submitted (SEP-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AE003698; AA014338.1; -
 SQ SEQUENCE 2347 AA; 257013 MM; 23BF5FC5FCACAE64 CRC64;
 Query Match 56.5%; Score 118; DB 5; Length 2347;
 Best Local Similarity 75.7%; Pred. No. 0.021;
 Matches 28; Conservative 3; Mismatches 6; Indels 0; Gaps 0;
 QY 3 AAAAAAAAAAAAAAAAAAKKKKKKKKKKK 39
 DB 1374 AAAAAAAAAAAAAAAAAEQAQKKNKKQAKK 1410
 RESULT 3
 Q9VG05 PRELIMINARY; PRT; 2451 AA.
 ID Q9VG05;
 AC Q9VG05;
 DT 01-MAY-2000 (TREMBLrel. 13, Created)
 DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
 DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
 DE CG7518 protein.
 GN CG7518.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydriidae; Drosophilidae; Drosophila.
 OC NCBI_taxid=7227;
 (1)
 RP SEQUENCE FROM N.A.
 RC STRAIN=BERKELEY;
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celniker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galie R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutcliffe G.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.H., Blazer R.G., Champe M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
 RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfankuch C., Baldwin D.,
 RA Bailew R.M., Basu A., Baxendale J., Bayraktaroglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Berman B.P., Bhandari D., Bolshakov S.,
 RA Bortova D., Botchan M.R., Bouck J., Brokstein P., Brothier P.,
 RA Burris K.C., Busan D.A., Butler H., Cadieu E., Center A., Chandra I.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,
 RA Durbin K.J., Evangelista C.C., Ferraz C., Ferreira S., Fleischmann W.,
 RA Foster C., Gabrielian A.E., Gary N.S., Gelbart W.M., Glaeser K.,
 RA Glodex A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
 RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
 RA Hostin D., Houston K.A., Howland T.J., Mei M.H., Ibegwan C.,
 RA Jatali M., Kalush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
 RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
 RA Laslo P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
 RA Liu X., Mattei B., McIntosh T.C., McLeod M.P., McPherson D.,

RA Merklov G., Milshina N.V., Mobarry C., Morris J., Moshrefi A.,
RA Mount S.M., Moy M., Murphy B., Murphy L., Wozny D.M., Nelson D.L.,
RA Nelson D.R., Nelson K.A., Nixon K., Nussekem D.R., Paclet J.M.,
RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Fui V., Reese M.G.,
RA Reinert K., Remington K., Saunders R.D.C., Scheller F., Shen H.,
RA Shie B.C., Siden-Klamos I., Simpson M., Skupski M.P., Smith T.,
RA Spier E., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Svirskaes R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Welzenbach J.,
RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
RT "The genome sequence of Drosophila melanogaster.";
RL Science 287:2185-2195(2000).
DR EMBL; AE003698; AAF54888.2; -.
DR FlyBase; FBgn0038108; CG7518.
DR InterPro; IPR001005; MYB_DNA_binding.
DR PROSITE; PS00037; MYB.1.1.
SQ SEQUENCE 2451 AA; 266959 MW; 088A2293F27481E2 CRC64;

Query Match 56.5%; Score 118; DB 5; Length 2451;
Best Local Similarity 75.7%; Pred. No. 0.022;
Matches 28; Conservative 3; Mismatches 6; Indels 0; Gaps 0;

OY 3 AAAAAAAAAAAAAAAAAAKKKKKKKKKKK 39
DB 1374 AAAAAAAAAAAAAAAAAEOKAKLKKKKKK 1410

RESULT 4

O8T2U7 PRELIMINARY; PRT; 791 AA.

AC O8T2U7; PRELIMINARY; PRT; 791 AA.
DT 01-JUN-2002 (TRENBLrel. 21, Created)
DT 01-JUN-2002 (TRENBLrel. 21, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE Hypothetical 92.4 kDa protein.
OS Dictyostelium discoideum (slime mold).
OC Eukaryota; Mycetozoa; Dictyostelida; Dictyostelium.
OX NCBI_TaxID=44689;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-AX4;
RA Giesecker G., Eichinger L., Szafranski K., Pachebat J., Dear P.,
RA Lehmann R., Baumgart C., Parra G., April J.F., Gulgo R., Kumpf K.,
RA Tungsang B., Cox B., Quail M.A., Platzner M., Rosenthal A., Noegel A.A.;
RL "Sequence and Analysis of Chromosome 2 of Dictyostelium";
RT Submitted (MAR-2002) to the EMBL/Genbank/DBJ databases.
RU EMBL; AC115574; AAL92183.1; -.
DR InterPro; IPR005033; YEATS.
DR InterPro; IPR007087; Znf_C2H2.
DR Pfam; PF00366; YEATS.1.
DR SMART; SMO0355; ZNF_C2H2; 1.
KW Hypothetical protein.
SQ SEQUENCE 791 AA; 92375 MW; D6CCB6DEC92352C CRC64;

Query Match 51.2%; Score 107; DB 5; Length 791;
Best Local Similarity 56.8%; Pred. No. 0.07;
Matches 25; Conservative 0; Mismatches 19; Indels 0; Gaps 0;

OY 2 AAAAAAAAAAAAAAAAAAKKKKKKKKKKK 45
DB 746 EKEIETEMIGKEIEAEIEIEKKKKKKKKKKKKKKKKKK 789

RESULT 5

O3S807 PRELIMINARY; PRT; 129 AA.

AC O3S807; PRELIMINARY; PRT; 129 AA.
DT 01-JAN-1998 (TRENBLrel. 05, Created)
DT 01-JAN-1998 (TRENBLrel. 05, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)

DE MICROVASCULAR endothelial differentiation protein 2.
GN MD02.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC Tissue=Epithelium;
RX MEDLINE=96172708; PubMed=9511718;
RA Pirels F., Loser B., Marx M.;
RT "Differential expression of osteopontin, PC4, and CEC5, a novel mRNA
RT species, during in vitro angiogenesis.";
RL Exp. Cell Res. 239:1-10(1998).
DR EMBL; Y08769; CAA70022.1; -.
DR InterPro; IPR00719; Prot_kinase.
DR Pfam; PF00069; Kinase; 1.
DR ProDom; PD000001; Prot_kinase; 1.
DR PROSITE; PS00011; PROTEIN_KINASE_DOM; 1.
KW ATP-binding; Transferase.
SQ SEQUENCE 129 AA; 15080 MW; 38102272BBE2EDB4 CRC64;

Query Match 50.2%; Score 105; DB 11; Length 129;
Best Local Similarity 95.5%; Pred. No. 0.02;
Matches 21; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 24 AAKKKKKKKKKKKKKKKKKKK 45
DB 83 ASKKKKKKKKKKKKKKKKKKKK 104

RESULT 6

O9H6O7 PRELIMINARY; PRT; 720 AA.

AC O9H6O7; PRELIMINARY; PRT; 720 AA.
DT 01-MAR-2001 (TRENBLrel. 16, Created)
DT 01-MAR-2001 (TRENBLrel. 16, Last sequence update)
DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)
DE Hypothetical protein FLJ21979 (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Kawabata A., Hiki J., Kobatake N., Inagaki H., Ikema Y., Okamoto S.,
RA Okitani R., Ota T., Suzuki Y., Odayashi M., Nishi T., Shibahara T.,
RA Tanaka T., Nakamura Y., Isegaki T., Sugano S.;
RL "NEDO human cDNA sequencing project.";
RT Submitted (AUG-2000) to the EMBL/Genbank/DBJ databases.
RU EMBL; AK025632; BAB15196.1; -.
DR KW Hypothetical protein.
FT NON TER 720
SQ SEQUENCE 720 AA; 84029 MW; A6586FEAA953DOB CRC64;

Query Match 50.2%; Score 105; DB 4; Length 720;
Best Local Similarity 61.8%; Pred. No. 0.095;
Matches 21; Conservative 6; Mismatches 7; Indels 0; Gaps 0;

OY 12 AAAAAAAAAAAAAAAAAAKKKKKKKKKKK 45
DB 678 AKKSITNSDIVISIKKKKKKKKKKKKKKKKKKKKK 711

RESULT 7

O8CGI8 PRELIMINARY; PRT; 658 AA.

AC O8CGI8; PRELIMINARY; PRT; 658 AA.
DT 01-MAR-2003 (TRENBLrel. 23, Created)
DT 01-MAR-2003 (TRENBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)
DE Hypothetical protein (Fragment).
OS Mus musculus (Mouse).

```

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sclurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxId=10090.
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=FVB/N;
RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL, BC035210, AAH35210.1; -.
KW Hypothetical protein.
SQ SEQUENCE 658 AA; 73538 MW; 270E82146355FF9 CRC64;
FT NON TER 658
Query Match 49.8%; Score 104; DB 11; Length 658;
Best Local Similarity 64.7%; Pred. No. 0.11;
Matches 22; Conservative 3; Mismatches 9; Indels 0; Gaps 0.

Qy 11 AAAAAAAAAAAAAAAAAKKKKKKKKKKKKKKKKKK 44
Db 625 ATSTTCTATVQAASSKKKKKKKKKKKKKKKKKK 658

RESULT 8
Q8LOP6 PRELIMINARY; PRT; 113 AA.
ID Q8LOP6
AC Q8LOP6
DT 01-OCT-2002 (TREMBLrel. 22, Created)
DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE OJ117_G01.13 protein.
GN OJ117_G01.13
OS Oryza sativa (japonica cultivar-group).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzae; Oryza.
OX NCBI_TaxId=39947;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=cv. Nipponbare;
RC Sasaki T., Matsumoto T., Yamamoto K.;
RT "Oryza sativa nipponbare (GA3) genomic DNA, chromosome 1, BAC
clone:OJ117_G01."
RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL, AP003374, BAB93330.1; -.
SQ SEQUENCE 113 AA; 13660 MW; 597DB0DEB2AA3EF CRC64;

Query Match 49.3%; Score 103; DB 10; Length 113;
Best Local Similarity 73.3%; Pred. No. 0.027; 7; Indels 0; Gaps 0.
Matches 22; Conservative 1; Mismatches 7; Indels 0; Gaps 0.

Qy 16 AAAAAAAAAAAAAAAAAKKKKKKKKKKKKKKKKKK 45
Db 3 ATSLHHKKKKKKKKKKKKKKKKKKKKKKKKKK 32

RESULT 9
Q9LL82 PRELIMINARY; PRT; 467 AA.
ID Q9LL82
AC Q9LL82
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
DE Cbf5p.
OS Euglena gracilis.
OC Eukaryota; Euklenozoa; Euglenida; Euglenales; Euglena.
OX NCBI_TaxId=3039;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=streptomycin-bleached strain;
RX MEDLINE=20330353; PubMed=10871366;
RA Watanabe Y., Gray M.W.;
RT "Evolutionary appearance of genes encoding proteins associated with

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RT box H/ACA snRNAs: Cbf5p in Engelma girecillii, an early diverging
RT eukaryote, and candidate Garp and Nop10p homologs in
RT archaeobacteria."
RL Nucleic Acids Res. 28:2342-2352(2000).
DR EMBL; AF234319; AAF77119.1; -.
DR InterPro; IPR004802; CBf5.
DR InterPro; IPR002478; PUA.
DR InterPro; IPR002501; TruB_N.
DR InterPro; IPR004521; Unchar_dom_2.
DR Pfam; PF01472; PUA; 1.
DR Pfam; PF01509; TruB_N; 1.
DR SMART; SM00359; PUA; 1.
DR TIGRFAMs; TIGR00425; CBP5; 1.
DR TIGRFAMs; TIGR00451; unchar_dom_2; 1.
SO SEQUENCE 467 AA; 52385 MW; 741089B66507BA7B CRC64;

Query Match 48.8%; Score 102; DB 10; Length 467;
Best Local Similarity 55.8%; Pred. No. 0.12;
Matches 24; Conservative 5; Mismatches 14; Indels 0; Gaps 0;

OY 2 EAAAAAAAAAAAAAAAAAAAAAAAAKKKKKKKKKKKKKKKKKK 44
Db 420 EELVAAEAAKKREREAAAGDEKDAAKKAKKEKKKKKKKK 462

RESULT 10
O9LKR2 PRELIMINARY; PRT; 517 AA.
AC Q9LKR2;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Hypothetical 59.7 KDa protein.
GN TGN10 250.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eustosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxId=3702;
CK [1]
RN RP SEQUENCE FROM N.A.
RA D'Angelo M., Vezzi A., Modesto D., Pigazzi M., Valle G., Mewes H.W.,
RA Rued S., Lemke K., Mayer K.F.X., Queller F., Salanoubat M.,
RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
RN RP SEQUENCE FROM N.A.
RA EU Arabidopsis sequencing project;
RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AL353032; CAB88307.1; -.
DR InterPro; IPR001810; F-box.
DR InterPro; IPR005566; FBD.
DR Pfam; PF00646; F-box; 1.
DR Pfam; PF00579; FBD; 1.
DR SMART; SM00256; PBOX; 1.
DR PROSITE; PS50181; PBOX; 1.
KM Hypothetical protein.
SQ SEQUENCE 517 AA; 59689 MW; BC6D957D01F86770 CRC64;

Query Match 48.8%; Score 102; DB 10; Length 517;
Best Local Similarity 80.8%; Pred. No. 0.13;
Matches 21; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

OY 20 EAAAAAAAAAAAAAAAAAAAAAAAAKKKKKKKKKKKKKKKK 45
Db 440 ERVGGKKKKKKKKKKKKKKKKKKKKKKKKKKKK 465

RESULT 11
O14347 PRELIMINARY; PRT; 354 AA.
AC O14347;
DT 01-JUN-1998 (TrEMBLrel. 06, Created)
DT 01-JUN-1998 (TrEMBLrel. 06, Last sequence update)
```


Query Match	Best Local Similarity	Matches	Score	DB	Length	Indels	Gaps
01-JUN-2002 (TReMBLrel. 21, last sequence update)	49.1%;	1;	100.5;	10;	80;	11;	1;
01-OCT-2002 (TReMBLrel. 22, last annotation update)	49.1%;	1;	100.5;	10;	80;	11;	1;
Hypothetical 9.4 kDa protein.	49.1%;	1;	100.5;	10;	80;	11;	1;
OSUNBA0057L21.23.	49.1%;	1;	100.5;	10;	80;	11;	1;
Oryza sativa (Rice).	49.1%;	1;	100.5;	10;	80;	11;	1;
Oryza sativa (Rice).	49.1%;	1;	100.5;	10;	80;	11;	1;
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;	49.1%;	1;	100.5;	10;	80;	11;	1;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;	49.1%;	1;	100.5;	10;	80;	11;	1;
Ehretioideae; Oryzae; Oryza.	49.1%;	1;	100.5;	10;	80;	11;	1;
NCBI_TaxID=4530;	49.1%;	1;	100.5;	10;	80;	11;	1;
SEQUENCE FROM N.A.	49.1%;	1;	100.5;	10;	80;	11;	1;
STRAIN=cv. Nipponbare;	49.1%;	1;	100.5;	10;	80;	11;	1;
Buell C.R., Yuan Q., Ouyang S., Liu J., Mofatt K.S., Hill J.N.,	49.1%;	1;	100.5;	10;	80;	11;	1;
Gansberger K., Brenner M., Burgess S., Hance M., Shvartsbeyn M.,	49.1%;	1;	100.5;	10;	80;	11;	1;
Teltrin T., Riggs F., Hsiao J., Zilman V., Blum S., Pei G.,	49.1%;	1;	100.5;	10;	80;	11;	1;
Valankin S.E., Utceback T.R., Feldblum T.V., Kalb E., Quackenbush J.,	49.1%;	1;	100.5;	10;	80;	11;	1;
Salzberg S.L., White O., Fraser C.M.;	49.1%;	1;	100.5;	10;	80;	11;	1;
"Oryza sativa chromosome 10 BAC OSUNBA0057L21 genomic sequence."	49.1%;	1;	100.5;	10;	80;	11;	1;
Submitted (FEB-2002) to the EMBL/GenBank/DBJ databases.	49.1%;	1;	100.5;	10;	80;	11;	1;
EMBL; AC087589; AL179706.1; -.	49.1%;	1;	100.5;	10;	80;	11;	1;
Gravene; Q857D3; -.	49.1%;	1;	100.5;	10;	80;	11;	1;
Hypothetical protein.	49.1%;	1;	100.5;	10;	80;	11;	1;
SEQUENCE 80 AA; 9362 MW; 0177C86313B21D8 CRC64;	49.1%;	1;	100.5;	10;	80;	11;	1;
Query Match	Best Local Similarity	Matches	Score	DB	Length	Indels	Gaps
01-JUN-2002 (TReMBLrel. 21, last sequence update)	49.1%;	1;	100.5;	10;	80;	11;	1;
01-OCT-2002 (TReMBLrel. 22, last annotation update)	49.1%;	1;	100.5;	10;	80;	11;	1;
Hypothetical 9.4 kDa protein.	49.1%;	1;	100.5;	10;	80;	11;	1;
OSUNBA0057L21.23.	49.1%;	1;	100.5;	10;	80;	11;	1;
Oryza sativa (Rice).	49.1%;	1;	100.5;	10;	80;	11;	1;
Oryza sativa (Rice).	49.1%;	1;	100.5;	10;	80;	11;	1;
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;	49.1%;	1;	100.5;	10;	80;	11;	1;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;	49.1%;	1;	100.5;	10;	80;	11;	1;
Ehretioideae; Oryzae; Oryza.	49.1%;	1;	100.5;	10;	80;	11;	1;
NCBI_TaxID=4530;	49.1%;	1;	100.5;	10;	80;	11;	1;
SEQUENCE FROM N.A.	49.1%;	1;	100.5;	10;	80;	11;	1;
STRAIN=cv. Nipponbare;	49.1%;	1;	100.5;	10;	80;	11;	1;
Buell C.R., Yuan Q., Ouyang S., Liu J., Mofatt K.S., Hill J.N.,	49.1%;	1;	100.5;	10;	80;	11;	1;
Gansberger K., Brenner M., Burgess S., Hance M., Shvartsbeyn M.,	49.1%;	1;	100.5;	10;	80;	11;	1;
Teltrin T., Riggs F., Hsiao J., Zilman V., Blum S., Pei G.,	49.1%;	1;	100.5;	10;	80;	11;	1;
Valankin S.E., Utceback T.R., Feldblum T.V., Kalb E., Quackenbush J.,	49.1%;	1;	100.5;	10;	80;	11;	1;
Salzberg S.L., White O., Fraser C.M.;	49.1%;	1;	100.5;	10;	80;	11;	1;
"Oryza sativa chromosome 10 BAC OSUNBA0057L21 genomic sequence."	49.1%;	1;	100.5;	10;	80;	11;	1;
Submitted (FEB-2002) to the EMBL/GenBank/DBJ databases.	49.1%;	1;	100.5;	10;	80;	11;	1;
EMBL; AC087589; AL179706.1; -.	49.1%;	1;	100.5;	10;	80;	11;	1;
Gravene; Q857D3; -.	49.1%;	1;	100.5;	10;	80;		

OC Sordariaceae; Sordariaceae; Neurospora.
 OX NCBI_TaxID=5141;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Schulte U., Aign V., Honeisel J., Brandt P., Fartmann B., Holland R.,
 RA Nyakatura G., Mewes H.W., Mannhaupt G.;
 RL Submitted (May-2000) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA German Neurospora genome project;
 RL Submitted (Oct-2001) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AL356815; CAB92638.2; -.
 KW Hypothetical protein.
 SQ SEQUENCE 128 AA; 15157 MW; 8C7C65C3DFB70765 CRC64;

Query Match 47.8%; Score 100; DB 3; Length 128;
 Best Local Similarity 100.0%; Pred. No. 0.055;
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 26 KKKKKKKKKKKKKKKKKKK 45
 |||||
 Db 71 KKKKKKKKKKKKKKKKKKK 90

Search completed: January 30, 2004, 00:24:41
 Job time : 46.4789 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 30, 2004, 00:13:12 ; Search time 18.7981 Seconds
(without alignments)
225.098 Million cell updates/sec

Title: US-09-461-684C-5

Perfect score: 243
Sequence: 1 GLEFGAIGFIENGWEGMID.....KKKKKKKKKKKKKKKKKK 44

Scoring table:
BLOSUM62
Gapop 10.0 , Gapept 0.5

Searched: 283308 seqs, 9618682 residues

Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database : PIR_76:*
1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	134	55.1	550	1 HMTV82	hemagglutinin prec
2	134	55.1	550	1 HMTV83	hemagglutinin prec
3	134	55.1	550	1 HMTV77	hemagglutinin prec
4	134	55.1	550	1 HMTV80	hemagglutinin prec
5	134	55.1	550	1 HMTV33	hemagglutinin prec
6	134	55.1	550	1 HMTV89	hemagglutinin prec
7	134	55.1	550	1 HMTV21	hemagglutinin prec
8	134	55.1	550	1 HMTV98	hemagglutinin prec
9	134	55.1	550	1 HMTV15	hemagglutinin prec
10	134	55.1	550	2 JQ1153	hemagglutinin prec
11	134	55.1	550	2 JQ1154	hemagglutinin prec
12	134	55.1	550	2 JQ1155	hemagglutinin prec
13	134	55.1	566	1 HMTV8	hemagglutinin prec
14	134	55.1	566	1 HMTV84	hemagglutinin prec
15	134	55.1	566	1 HMTV84	hemagglutinin prec
16	134	55.1	566	1 HMTV84	hemagglutinin prec
17	134	55.1	570	1 A45591	hemagglutinin prec
18	134	55.1	570	2 S22013	hemagglutinin prec
19	134	55.1	561	1 HMTV49	hemagglutinin prec
20	133	54.7	561	1 HMTV84	hemagglutinin prec
21	132	54.3	565	1 HMTV81	hemagglutinin prec
22	132	54.3	565	1 HMTV81	hemagglutinin prec
23	132	54.3	566	1 HMTV6	hemagglutinin prec
24	132	54.3	567	1 HMTV6	hemagglutinin prec
25	131	53.9	362	2 S38637	hemagglutinin - in
26	131	53.9	550	1 HMTV86	hemagglutinin prec
27	131	53.9	560	1 HMTV86	hemagglutinin prec
28	131	53.9	565	1 HMTV82	hemagglutinin prec
29	131	53.9	565	1 HMTV84	hemagglutinin prec

30	131	53.9	565	1 HMTV85	hemagglutinin prec
31	131	53.9	565	1 HMTV86	hemagglutinin prec
32	131	53.9	565	1 HMTV87	hemagglutinin prec
33	131	53.9	565	1 HMTV88	hemagglutinin prec
34	131	53.9	565	1 HMTV89	hemagglutinin prec
35	131	53.9	565	1 HMTV89	hemagglutinin prec
36	131	53.9	565	1 HMTV89	hemagglutinin prec
37	131	53.9	565	2 S33703	hemagglutinin - in
38	131	53.9	570	2 S22014	hemagglutinin prec
39	131	53.9	570	2 S22015	hemagglutinin prec
40	131	53.9	570	2 S22016	hemagglutinin prec
41	131	53.9	570	2 S22017	hemagglutinin prec
42	131	53.9	570	2 S22018	hemagglutinin prec
43	131	53.9	570	2 S22020	hemagglutinin prec
44	131	53.9	570	2 S22021	hemagglutinin prec
45	131	53.9	570	2 S22029	hemagglutinin prec

ALIGNMENTS

```
RESULT 1
HMTV82      hemagglutinin precursor - influenza A virus (strain A/swine/126/82) (fragment)
C:Species:  influenza A virus
C>Date: 31-Mar-1989 #sequence_revision 31-Mar-1989 #text_change 18-Sep-1998
C:Accession: A29971
R:Kida, H.; Shortridge, K.F.; Webster, R.G.
Virology 162, 160-166, 1988
A:Title: Origin of the hemagglutinin gene of H3N2 influenza viruses from pigs in China.
A:Reference number: A94370; MUID:88101364; PMID:3336940
A:Accession: A29971
A:Molecule type: genomic RNA
A:Residues: 1-550 <KID>
A:Cross-references: GB:M19056; NID:g324208
A:Note: the sequence in Genbank entry FLAHABP, release 106, (PID:g324209) differs from tl
C:Genetics:
A:Map position: segment 4
C:Superfamily: influenza virus hemagglutinin
C:Keywords: glycoprotein, hemagglutinin; homotrimer; lipoprotein; thiolester bond
F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>
F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>
F:520-536/Domain: transmembrane #status predicted <TM1>
F:8,22,38,165,285,483/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:14-486,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted
F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match      55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 3.1e-06;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy      2 GLEFGAIGFIENGWEGMIDGMYG 24
Db      330 GLEFGAIGFIENGWEGMIDGMYG 352

RESULT 2
HMTV83      hemagglutinin precursor - influenza A virus (strain A/swine/81/78) (fragment)
C:Species:  influenza A virus
C>Date: 31-Mar-1989 #sequence_revision 31-Mar-1989 #text_change 18-Sep-1998
C:Accession: B29971
R:Kida, H.; Shortridge, K.F.; Webster, R.G.
Virology 162, 160-166, 1988
A:Title: Origin of the hemagglutinin gene of H3N2 influenza viruses from pigs in China.
A:Reference number: A94370; MUID:88101364; PMID:3336940
A:Accession: B29971
A:Molecule type: genomic RNA
A:Residues: 1-550 <KID>
A:Cross-references: GB:M19057; NID:g324210
A:Note: the sequence in Genbank entry FLAHABP, release 106, (PID:g324211) differs from tl
C:Genetics:
A:Map position: segment 4
```

C:Superfamily: influenza virus hemagglutinin
C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond
F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>
F:300-550/Product: hemagglutinin HA2 #status predicted <HA2>
F:520-536/Domains: transmembrane #status predicted <TM1>
F:8,22,38,165,285,483/Binding site: carbohydrate (asn) (covalent) #status predicted
F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted
F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 3.1e-06;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMIDGWTG 24
DB 330 GLFGAIAGFIENGWEGMIDGWTG 352

RESULT 3
HMI177
hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/5/77) (fragment)
N:Contains: hemagglutinin HA1; hemagglutinin HA2
C:Species: influenza A virus
C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 16-Jul-1999
C:Accession: A27813
R:Kida, H.; Kawaka, Y.; Naeve, C.W.; Webster, R.G.
Virology 159, 109-119, 1987
A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.
A:Reference number: A94363; PMID:87265458; PMID:2440178
A:Accession: A27813
A:Molecule type: genomic RNA
A:Residues: 1-550 <KID>
A:Cross-references: GB:M16737; NID:G324081; PIDN:AAA43143.1; PID:G324082

C:Genetics:
A:Map position: segment 4
C:Superfamily: influenza virus hemagglutinin
C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond
F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>
F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>
F:520-536/Domains: transmembrane #status predicted <TM1>
F:8,22,38,165,285,483/Binding site: carbohydrate (asn) (covalent) #status predicted
F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted
F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 3.1e-06;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMIDGWTG 24
DB 330 GLFGAIAGFIENGWEGMIDGWTG 352

RESULT 4
HMI180
hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/8/80) (fragment)
N:Contains: hemagglutinin HA1; hemagglutinin HA2
C:Species: influenza A virus
C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 18-Sep-1998
C:Accession: B27813
R:Kida, H.; Kawaka, Y.; Naeve, C.W.; Webster, R.G.
Virology 159, 109-119, 1987
A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.
A:Reference number: A94363; PMID:87265458; PMID:2440178
A:Accession: B27813
A:Molecule type: genomic RNA
A:Residues: 1-550 <KID>
A:Cross-references: GB:M16738; NID:G324083

A:Note: the translation in Fig. 2 is inconsistent with the nucleotide sequence in Fig. 1
C:Genetics:
A:Map position: segment 4
C:Superfamily: influenza virus hemagglutinin
C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>
F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>
F:520-536/Domains: transmembrane #status predicted <TM1>
F:8,22,38,165,285,483/Binding site: carbohydrate (asn) (covalent) #status predicted
F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted
F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 3.1e-06;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMIDGWTG 24
DB 330 GLFGAIAGFIENGWEGMIDGWTG 352

RESULT 5
HMI173
hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/33/80) (fragment)
N:Contains: hemagglutinin HA1; hemagglutinin HA2
C:Species: influenza A virus
C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 16-Jul-1999
C:Accession: C27813
R:Kida, H.; Kawaka, Y.; Naeve, C.W.; Webster, R.G.
Virology 159, 109-119, 1987
A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.
A:Reference number: A94363; PMID:87265458; PMID:2440178
A:Accession: C27813
A:Molecule type: genomic RNA
A:Residues: 1-550 <KID>
A:Cross-references: GB:M16739; NID:G324085; PIDN:AAA43145.1; PID:G324086

C:Genetics:
A:Map position: segment 4
C:Superfamily: influenza virus hemagglutinin
C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond
F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>
F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>
F:520-536/Domains: transmembrane #status predicted <TM1>
F:8,22,38,165,285,483/Binding site: carbohydrate (asn) (covalent) #status predicted
F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted
F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 3.1e-06;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMIDGWTG 24
DB 330 GLFGAIAGFIENGWEGMIDGWTG 352

RESULT 6
HMI189
hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/7/82) (fragment)
N:Contains: hemagglutinin HA1; hemagglutinin HA2
C:Species: influenza A virus
C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 16-Jul-1999
C:Accession: D27813
R:Kida, H.; Kawaka, Y.; Naeve, C.W.; Webster, R.G.
Virology 159, 109-119, 1987
A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.
A:Reference number: A94363; PMID:87265458; PMID:2440178
A:Accession: D27813
A:Molecule type: genomic RNA
A:Residues: 1-550 <KID>
A:Cross-references: GB:M16740; NID:G324087; PIDN:AAA43146.1; PID:G324088

A:Note: the translation in Fig. 2 is inconsistent with the nucleotide sequence in Fig. 1
C:Genetics:
A:Map position: segment 4
C:Superfamily: influenza virus hemagglutinin
C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond
F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>
F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>
F:520-536/Domains: transmembrane #status predicted <TM1>

F:8,22,38,165,285,483/Binding site: carboxylate (Asn) (covalent) #status predicted
 F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted
 F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 55.1%; Score 134; DB 1; Length 550;
 Best Local Similarity 100.0%; Pred. No. 3.1e-06;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

2 GLFGAIAGFIENGWEGMIDGWYG 24
 Db 330 GLFGAIAGFIENGWEGMIDGWYG 352

RESULT 7

HMIIV21

hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/21/82) (fragment)
 N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: influenza A virus
 C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 18-Sep-1998
 C:Accession: E27813

R:Kida, H.; Kawoka, Y.; Naeve, C.W.; Webster, R.G.

Virology 159, 109-119, 1987

A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.

A:Reference number: A94363; MUID:87265458; PMID:2440178

A:Accession: E27813

A:Molecule type: genomic RNA

A:Residues: 1-550 <KID>

A:Cross-references: GB:M16741; NID:9324089

C:Genetics:

A:Map position: segment 4

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>

F:520-536/Domains: transmembrane #status predicted <TM1>

F:7,22,38,165,285,483/Binding site: carboxylate (Asn) (covalent) #status predicted

F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted

Query Match 55.1%; Score 134; DB 1; Length 550;
 Best Local Similarity 100.0%; Pred. No. 3.1e-06;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

2 GLFGAIAGFIENGWEGMIDGWYG 24
 Db 330 GLFGAIAGFIENGWEGMIDGWYG 352

RESULT 8

HMIIV98

hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/9/85) (fragment)
 N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: influenza A virus

C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 18-Sep-1998
 C:Accession: F27813

R:Kida, H.; Kawoka, Y.; Naeve, C.W.; Webster, R.G.

Virology 159, 109-119, 1987

A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.

A:Reference number: A94363; MUID:87265458; PMID:2440178

A:Accession: F27813

A:Molecule type: genomic RNA

A:Residues: 1-550 <KID>

A:Cross-references: GB:M16742; NID:9324091

C:Genetics:

A:Map position: segment 4

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>

F:520-536/Domains: transmembrane #status predicted <TM1>

F:8,22,38,165,285,483/Binding site: carboxylate (Asn) (covalent) #status predicted

F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted

F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 55.1%; Score 134; DB 1; Length 550;
 Best Local Similarity 100.0%; Pred. No. 3.1e-06;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

2 GLFGAIAGFIENGWEGMIDGWYG 24
 Db 330 GLFGAIAGFIENGWEGMIDGWYG 352

RESULT 9

HMIIV15

hemagglutinin precursor - influenza A virus (strain A/duck/Hokkaido/10/85) (fragment)
 N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: influenza A virus

C:Date: 30-Jun-1989 #sequence_revision 30-Jun-1989 #text_change 16-Jul-1999
 C:Accession: G27813

R:Kida, H.; Kawoka, Y.; Naeve, C.W.; Webster, R.G.

Virology 159, 109-119, 1987

A:Title: Antigenic and genetic conservation of H3 influenza virus in wild ducks.

A:Reference number: A94363; MUID:87265458; PMID:2440178

A:Accession: G27813

A:Molecule type: genomic RNA

A:Residues: 1-550 <KID>

A:Cross-references: GB:M16743; NID:9324093; PIDN:AAA3149.1; PID:9324094

C:Genetics:

A:Map position: segment 4

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; hemagglutinin; homotrimer; lipoprotein; thiolester bond

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-550/Product: hemagglutinin HA2 #status predicted <HA2>

F:520-536/Domains: transmembrane #status predicted <TM1>

F:8,22,38,165,285,483/Binding site: carboxylate (Asn) (covalent) #status predicted

F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted

F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 55.1%; Score 134; DB 1; Length 550;
 Best Local Similarity 100.0%; Pred. No. 3.1e-06;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

2 GLFGAIAGFIENGWEGMIDGWYG 24
 Db 330 GLFGAIAGFIENGWEGMIDGWYG 352

RESULT 10

JQ1153

hemagglutinin precursor - influenza A virus (strain A/duck/Hong Kong/7/75) (fragment)
 N:Contains: hemagglutinin HA1; hemagglutinin HA2

C:Species: influenza A virus

C:Date: 31-Mar-1992 #sequence_revision 31-Mar-1992 #text_change 20-Jun-2000
 C:Accession: JQ1153

R:Yaounda, J.; Shortridge, K.F.; Shimizu, Y.; Kida, H.

J. Gen. Virol. 72, 2007-2010, 1991

A:Title: Molecular evidence for a role of domestic ducks in the introduction of avian H3

A:Reference number: JQ1153; MUID:91341491; PMID:1875195

A:Accession: JQ1153

A:Molecule type: genomic RNA

A:Residues: 1-550 <YAS>

A:Cross-references: GB:D00929; NID:9221279; PIDN:BA00769.1; PID:9221280

A:Note: the authors translated the codon GGG for residue 218 as Glu

A:Note: residues 528-532 are not shown in this publication

C:Superfamily: influenza virus hemagglutinin

C:Keywords: glycoprotein; homotrimer

F:1-328/Product: hemagglutinin HA1 #status predicted <HA1>

F:330-545/Product: hemagglutinin HA2 #status predicted <HA2>

F:8,22,38,165,285,483/Binding site: carboxylate (Asn) (covalent) #status predicted

F:14-466,52-277,64-76,139-473,281-305/Disulfide bonds: #status predicted

F:539,546,549/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 55.1%; Score 134; DB 2; Length 550;
 Best Local Similarity 100.0%; Pred. No. 3.1e-06;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

2 GLFGAIAGFIENGWEGMIDGWYG 24

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Db      330 GLFGAIAAGFIENGWEGMIDGWYG 352

RESULT 11
JQ1154 hemagglutinin precursor - influenza A virus (strain A/goose/Hong Kong/10/76) (fragment)
N:Contains: hemagglutinin HA1, hemagglutinin HA2
C:Species: influenza A virus
C>Date: 31-Mar-1992 #sequence_revision 31-Mar-1992 #text_change 20-Jun-2000
C:Accession: JQ1154
R:Yaounda, J.; Shortridge, K.F.; Shmizu, Y.; Kida, H.
J. Gen. Virol. 72, 2007-2010, 1991
A>Title: Molecular evidence for a role of domestic ducks in the introduction of avian H3
A:Reference number: JQ1153; MUID:91341491; PMID:1875195
A:Accession: JQ1154
A:Molecule type: genomic RNA
A:Residues: 1-550 <YAS>
A:Cross-references: GB:D00930; NID:9221273; PIDN:BA00770.1; PID:9221274
A>Note: the authors translated the codon GGG for residue 218 as G1U
A>Note: residues 528-532 are not shown in this publication
C:Superfamily: influenza virus hemagglutinin
C:Keywords: glycoprotein; homotrimer
F:1-358/Product: hemagglutinin HA1 #status predicted <HA1>
F:330-545/Product: hemagglutinin HA2 #status predicted <HA2>
F:8,22,38,165,285,483/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match      55.1%; Score 134; DB 2; Length 550;
Best Local Similarity 100.0%; Pred. No. 3.1e-06;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      2 GLFGAIAAGFIENGWEGMIDGWYG 24
Db      330 GLFGAIAAGFIENGWEGMIDGWYG 352

RESULT 12
JQ1155 hemagglutinin precursor - influenza A virus (strain A/duck/Hong Kong/64/76) (fragment)
N:Contains: hemagglutinin HA1, hemagglutinin HA2
C:Species: influenza A virus
C>Date: 31-Mar-1992 #sequence_revision 31-Mar-1992 #text_change 20-Jun-2000
C:Accession: JQ1155
R:Yaounda, J.; Shortridge, K.F.; Shmizu, Y.; Kida, H.
J. Gen. Virol. 72, 2007-2010, 1991
A>Title: Molecular evidence for a role of domestic ducks in the introduction of avian H3
A:Reference number: JQ1153; MUID:91341491; PMID:1875195
A:Accession: JQ1155
A:Molecule type: genomic RNA
A:Residues: 1-550 <YAS>
A:Cross-references: GB:D00931; NID:9221277; PIDN:BA00771.1; PID:9221278
A>Note: the authors translated the codon GGG for residue 218 as G1U, GCC for residue 538
A>Note: residues 528-532 are not shown in this publication
C:Superfamily: influenza virus hemagglutinin
C:Keywords: glycoprotein; homotrimer
F:1-358/Product: hemagglutinin HA1 #status predicted <HA1>
F:330-545/Product: hemagglutinin HA2 #status predicted <HA2>
F:8,22,38,165,285,483/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match      55.1%; Score 134; DB 2; Length 550;
Best Local Similarity 100.0%; Pred. No. 3.1e-06;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      2 GLFGAIAAGFIENGWEGMIDGWYG 24
Db      330 GLFGAIAAGFIENGWEGMIDGWYG 352

RESULT 13
HMI1VH hemagglutinin precursor - influenza A virus
C:Species: influenza A virus
C>Date: 28-Feb-1981 #sequence_revision 28-Feb-1981 #text_change 22-Oct-1999
C:Accession: A93705; A93233; A04051; A93231; A94441

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R:Both, G.W.; Sleight, M.J.
Nucleic Acids Res. 8, 2561-2575, 1980
A>Title: Complete nucleotide sequence of the haemagglutinin gene from a human influenza A
A:Reference number: A93705; MUID:81053698; PMID:6253883
A:Accession: A93705
A:Molecule type: Genomic RNA
A:Residues: 1-566 <BOT>
A:Cross-references: GB:V01103
A:Experimental source: strain A/NT/60/68/29C
A>Note: Human influenza strain A/NT/60/68/29C is a laboratory-isolated variant of A/NT/60
R:Dopheide, T.A.; Ward, C.W.
FBS Lett. 110, 181-183, 1980
A>Title: The disulphide bonds of a Hong Kong influenza virus hemagglutinin.
A:Reference number: A91276; MUID:80179105; PMID:6768586
A:Contents: annotation; disulfide bonds
A:Getting, M.J.; Bye, U.; Skehel, J.; Waterfield, M.
Nature 287, 301-306, 1980
A>Title: Cloning and DNA sequence of double-stranded copies of haemagglutinin genes from
A:Reference number: A93233; MUID:81030852; PMID:7421990
A:Accession: A93233
A:Molecule type: genomic RNA
A:Residues: 1-24, 'S', '26', 'D', '28-159', 'G', '161-197', 'I', '199-241', 'L', '243-249' <GRT>
A:Experimental source: strain X-31[H3]
C:Superfamily: influenza virus hemagglutinin
C:Keywords: hemagglutinin; homotrimer; lipoprotein; thiolester bond
F:1-16/Domain: signal sequence #status predicted <SIG>
F:17-244/Product: hemagglutinin HA1 #status predicted <HA1>
F:346-566/Product: hemagglutinin HA2 #status predicted <HA2>
F:536-552/Domain: transmembrane #status predicted <TM1>
F:30-482,82,92,155-489,297-321/Disulfide bonds: #status experimental
F:555,562,565/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match      55.1%; Score 134; DB 1; Length 566;
Best Local Similarity 100.0%; Pred. No. 3.2e-06;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      2 GLFGAIAAGFIENGWEGMIDGWYG 24
Db      346 GLFGAIAAGFIENGWEGMIDGWYG 368

RESULT 14
HMI1VH hemagglutinin precursor - influenza A virus (strain A/ichi/2/68)
N:Contains: hemagglutinin HA1, hemagglutinin HA2
C:Species: influenza A virus
C>Date: 19-May-1994 #sequence_revision 19-May-1994 #text_change 16-Jul-1999
C:Accession: A93231; A04051
R:Verhoeven, M.; Fang, R.; Man Jou, W.; Devos, R.; Huybreoek, D.; Saman, E.; Fiers, W.
Nature 286, 771-776, 1980
A>Title: Antigenic drift between the haemagglutinin of the Hong Kong influenza strains A
A:Reference number: A93231; MUID:80254693; PMID:7402351
A:Accession: A93231
A:Molecule type: Genomic RNA
A:Residues: 1-566 <VER>
A:Cross-references: GB:J02090; NID:9324131; PIDN:AA43178.1; PID:9324132
C:Superfamily: influenza virus hemagglutinin
C:Keywords: hemagglutinin; homotrimer; lipoprotein; thiolester bond
F:1-16/Domain: signal sequence #status predicted <SIG>
F:17-244/Product: hemagglutinin HA1 #status predicted <HA1>
F:346-566/Product: hemagglutinin HA2 #status predicted <HA2>
F:555,562,565/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match      55.1%; Score 134; DB 1; Length 566;
Best Local Similarity 100.0%; Pred. No. 3.2e-06;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY      2 GLFGAIAAGFIENGWEGMIDGWYG 24
Db      346 GLFGAIAAGFIENGWEGMIDGWYG 368

RESULT 15

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HMI17M
 hemagglutinin precursor - influenza A virus (strain A/Mem/102/72)
 N:Contains: hemagglutinin HA1; hemagglutinin HA2
 C:Species: influenza A virus
 C:Date: 19-May-1994 #sequence_revision 19-May-1994 #text_change 31-Mar-2000
 C:Accession: A94441; A04051
 R:Stleigh, M.J.; Both, G.W.; Brownlee, G.G.; Bender, V.J.; Moss, B.A.
 in Structure and Variation in Influenza Virus, Laver, G., eds., pp.69-79, F
 A:Title: The haemagglutinin gene of influenza A virus: nucleotide sequence analysis of C
 A:Reference number: A94441
 A:Accession: A94441
 A:Molecule type: genomic RNA
 A:Residues: 1-566 <SLR>
 C:Superfamily: influenza virus hemagglutinin
 C:Keywords: hemagglutinin; homotrimer; lipoprotein; thiolester bond
 F:1-16/Domain: signal sequence #status predicted <SIG>
 F:17-344/Product: hemagglutinin HA1 #status predicted <HA1>
 F:346-566/Product: hemagglutinin HA2 #status predicted <HA2>
 F:555,562,565/Binding site: palmitate (Cys) (covalent) #status predicted

Query Match 55.1%; Score 134; DB 1; Length 566;
 Best Local Similarity 100.0%; Pred. No. 3.2e-06;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIGFIENGWEGMIDGWYG 24
 |||||
 DB 346 GLFGAIGFIENGWEGMIDGWYG 368

Search completed: January 30, 2004, 00:26:22
 Job time : 18.7981 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 29, 2004, 23:58:27 ; Search time 9.91549 Seconds
(without alignments)
208.681 Million cell updates/sec

Title: US-09-461-684C-5
Perfect score: 243
Sequence: 1 CGEFGAAGTENGWEGMID.....KKKKKKKKKKKKKKKKKK 44

Scoring table: BIOSUM62
Gapop 10.0, Gapext 0.5

Searched: 127863 seqs, 47026705 residues
Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database: SwissProt_41.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	134	55.1	550	1 HEMA_IADH1	P12582 Influenza a
2	134	55.1	550	1 HEMA_IADH2	P12583 Influenza a
3	134	55.1	550	1 HEMA_IADH3	P12584 Influenza a
4	134	55.1	550	1 HEMA_IADH4	P12585 Influenza a
5	134	55.1	550	1 HEMA_IADH5	P12586 Influenza a
6	134	55.1	550	1 HEMA_IADH6	P12587 Influenza a
7	134	55.1	550	1 HEMA_IADH7	P12588 Influenza a
8	134	55.1	550	1 HEMA_IADH8	P43257 Influenza a
9	134	55.1	550	1 HEMA_IADH9	P43258 Influenza a
10	134	55.1	550	1 HEMA_IADH10	P43260 Influenza a
11	134	55.1	550	1 HEMA_IADH11	P11133 Influenza a
12	134	55.1	550	1 HEMA_IADH12	P11134 Influenza a
13	134	55.1	550	1 HEMA_IADH13	P03437 Influenza a
14	134	55.1	550	1 HEMA_IADH14	P03438 Influenza a
15	134	55.1	550	1 HEMA_IADH15	P03442 Influenza a
16	134	55.1	550	1 HEMA_IADH16	P26134 Influenza a
17	134	55.1	550	1 HEMA_IADH17	P26135 Influenza a
18	134	55.1	550	1 HEMA_IADH18	P03439 Influenza a
19	134	55.1	550	1 HEMA_IADH19	P03436 Influenza a
20	134	55.1	550	1 HEMA_IADH20	P26094 Influenza a
21	134	55.1	550	1 HEMA_IADH21	P26101 Influenza a
22	133	54.7	561	1 HEMA_IADH22	P12581 Influenza a
23	133	54.7	561	1 HEMA_IADH23	P12582 Influenza a
24	132	54.3	565	1 HEMA_IADH24	P17000 Influenza a
25	132	54.3	565	1 HEMA_IADH25	P17002 Influenza a
26	132	54.3	566	1 HEMA_IADH26	P03440 Influenza a
27	132	54.3	566	1 HEMA_IADH27	P26139 Influenza a
28	132	54.3	567	1 HEMA_IADH28	P03435 Influenza a
29	131	53.9	560	1 HEMA_IADH29	P12589 Influenza a
30	131	53.9	560	1 HEMA_IADH30	P03458 Influenza a
31	131	53.9	565	1 HEMA_IADH31	P16994 Influenza a
32	131	53.9	565	1 HEMA_IADH32	P16995 Influenza a
33	131	53.9	565	1 HEMA_IADH33	P19699 Influenza a

34	131	53.9	565	1 HEMA_IADH34	P16996 Influenza a
35	131	53.9	565	1 HEMA_IADH35	P16997 Influenza a
36	131	53.9	565	1 HEMA_IADH36	P16998 Influenza a
37	131	53.9	565	1 HEMA_IADH37	P16999 Influenza a
38	131	53.9	565	1 HEMA_IADH38	P16999 Influenza a
39	131	53.9	565	1 HEMA_IADH39	P16999 Influenza a
40	131	53.9	565	1 HEMA_IADH40	P16999 Influenza a
41	131	53.9	565	1 HEMA_IADH41	P16999 Influenza a
42	131	53.9	565	1 HEMA_IADH42	P16999 Influenza a
43	131	53.9	565	1 HEMA_IADH43	P16999 Influenza a
44	131	53.9	565	1 HEMA_IADH44	P16999 Influenza a
45	131	53.9	565	1 HEMA_IADH45	P16999 Influenza a

ALIGNMENTS

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RESULT 1
HEMA_IADH1 STANDARD; PRT; 550 AA.
AC P12582; Q84021; Q84022;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-OCT-1989 (Rel. 12, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [contains: Hemagglutinin HA1 chain;
DE Hemagglutinin HA2 chain] (Fragment).
GN HA.
OS Influenza A virus (strain A/Duck/Hokkaido/5/77).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=11357;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87265458; PubMed=2440178;
RA Kida H., Kawasaka Y., Naeye C.W., Webster R.G.;
RT "Antigenic and genetic conservation of H3 influenza virus in wild
RT ducks.";
RL Virology 159:109-119 (1987).
CC -!- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC -!- CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -!- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -!- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
CC
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CC or send an email to license@ebi.ac.uk).
CC
CC EMBL; P12582; AAA43143.1; -.
CC DR HSSP; P03437; 3HWG.
CC DR InterPro; IPR001364; Hemagglutinin.
CC DR Pfam; PF00509; Hemagglutinin; 1.
CC DR PRINTS; PR00329; HEMAGGLUTININ2.
CC DR ProDom; PD000225; Hemagglutn; 1.
CC KW Envelope protein; Hemagglutinin; Glycoprotein.
CC FT NON TER 1
CC FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
CC FT CHAIN 330 550 HEMAGGLUTININ HA2 CHAIN.
CC FT CARBOHYD 8
CC FT CARBOHYD 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC FT CARBOHYD 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
CC SQ SEQUENCE 550 AA; 61705 MW; 7E7ACFE716FC969A CRC64;

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 2,2e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

OY 2 GLFGAIAFIENGMEGMDWG 24
DB 330 GLFGAIAFIENGMEGMDWG 352

RESULT 2

HEMA_IADH2 STANDARD: PRT: 550 AA.

AC P12583; Q84011; (Rel. 12, Created)
DT 01-OCT-1989 (Rel. 12, Last sequence update)
DT 01-APR-1990 (Rel. 14, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
Hemagglutinin HA2 chain] (Fragment).

GN HA.
OS Influenza A virus (strain A/Duck/Hokkaido/8/80).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=11358;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87265458; PubMed=2440178;
RA Kida H., Kawaoaka Y., Naeve C.W., Webster R.G.;
RT "Antigenic and genetic conservation of H3 influenza virus in wild
ducks";

RL Virology 159:109-119(1987).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
(HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.

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or send an email to license@isb-sib.ch).

CC EMBL; M16738; AAA43144.1; -
DR CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
DR PIR; B27813; HMI780.
DR HSSP; P03437; 2VIU.
DR InterPro: IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTIN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Hemagglutinin; Glycoprotein.

FT NON TER 1 1
FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CHAIN 330 550 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 8 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 137 137 K -> N (IN PIR DATA BANK).
SQ SEQUENCE 550 AA; 61659 MW; A107023ACC9CC35 CRC64;

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 2.2e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAIAFIENGMEGMDWG 24
DB 330 GLFGAIAFIENGMEGMDWG 352

RESULT 3
HEMA_IADH3 STANDARD: PRT: 550 AA.

AC P12584; Q84012; Q89793;

DT 01-OCT-1989 (Rel. 12, Created)

DT 01-OCT-1989 (Rel. 12, Last sequence update)

DT 16-OCT-2001 (Rel. 40, Last annotation update)

DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
Hemagglutinin HA2 chain] (Fragment).

GN HA.
OS Influenza A virus (strain A/Duck/Hokkaido/33/80).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=11359;

RP SEQUENCE FROM N.A.
RX MEDLINE=87265458; PubMed=2440178;
RA Kida H., Kawaoaka Y., Naeve C.W., Webster R.G.;
RT "Antigenic and genetic conservation of H3 influenza virus in wild
ducks";

RL Virology 159:109-119(1987).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
(HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.

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or send an email to license@isb-sib.ch).

CC EMBL; M16739; AAA43145.1; -
DR CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
DR HSSP; P03437; 2VIU.
DR InterPro: IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTIN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Hemagglutinin; Glycoprotein.

FT NON TER 1 1
FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CHAIN 330 550 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 8 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 550 AA; 61577 MW; 6C30BF67CFDCB7DE CRC64;

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 2.2e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAIAFIENGMEGMDWG 24
DB 330 GLFGAIAFIENGMEGMDWG 352

RESULT 4
HEMA_IADH4 STANDARD: PRT: 550 AA.

AC P12585; Q84013; Q84014;

DT 01-OCT-1989 (Rel. 12, Created)

DT 01-OCT-1989 (Rel. 12, Last sequence update)

DT 16-OCT-2001 (Rel. 40, Last annotation update)

DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
Hemagglutinin HA2 chain] (Fragment).

GN HA.
OS Influenza A virus (strain A/Duck/Hokkaido/7/82).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=11360;

RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87265458; PubMed=2440178;
 RA Kida H., Kawaka Y., Nave C.W., Webster R.G.;
 RT "Antigenic and genetic conservation of H3 influenza virus in wild
 ducks.";
 RL Virology 159:109-119(1987).
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION.
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 CC -----
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 CC -----
 CC EMBL; M16740; AAA43146.1; -.
 CC DR HSSP; P03437; 2V1U.
 CC DR InterPro; IPR001364; Hemagglutn.
 CC DR Pfam; PF00509; Hemagglutinin; 1.
 CC DR PRINTS; PR00329; HEMAGGLUTN12.
 CC DR ProDom; PD000225; Hemagglutn; 1.
 CC KW Envelope protein; Hemagglutinin; Glycoprotein.
 CC FT NON TER 1 328
 CC FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
 CC FT CARBOHYD 8 550 HEMAGGLUTININ HA2 CHAIN.
 CC FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC SQ SEQUENCE 550 AA; 61664 MW; A16B2CF8CBBDB9D0 CRC64;
 CC
 CC Query Match 55.1%; Score 134; DB 1; Length 550;
 CC Best Local Similarity 100.0%; Pred. No. 2.2e-07;
 CC Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 CC
 CC 2 GLFGAIGFIENGWEGMIDWYG 24
 CC DB 330 GLFGAIGFIENGWEGMIDWYG 352
 CC
 CC RESULT 5
 CC HEMA_IADH5 STANDARD; PRT; 550 AA.
 CC ID HEMA_IADH5 Q84015; Q84016;
 CC AC P12586; Q84015; Q84016;
 CC DT 01-OCT-1989 (Rel. 12, Created)
 CC DT 01-APR-1990 (Rel. 14, Last sequence update)
 CC DT 16-OCT-2001 (Rel. 40, Last annotation update)
 CC DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
 CC DE Hemagglutinin HA2 chain] (Fragment).
 CC GN HA.
 CC OS Influenza A virus (strain A/Duck/Hokkaido/21/82).
 CC OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 CC CC Influenza A viruses; Influenzavirus A.
 CC OX NCBI_TaxID=11361;
 CC RN [1]
 CC RP SEQUENCE FROM N.A.
 CC RX MEDLINE=87265458; PubMed=2440178;
 CC RA Kida H., Kawaka Y., Nave C.W., Webster R.G.;
 CC RT "Antigenic and genetic conservation of H3 influenza virus in wild
 CC ducks.";
 CC RL Virology 159:109-119(1987).
 CC CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION.
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.

CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
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 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL; M16741; AAA43147.1; -.
 CC DR PIR; E27813; HMTV21.
 CC DR HSSP; P03437; 2V1U.
 CC DR InterPro; IPR001364; Hemagglutn.
 CC DR Pfam; PF00509; Hemagglutinin; 1.
 CC DR PRINTS; PR00329; HEMAGGLUTN12.
 CC DR ProDom; PD000225; Hemagglutn; 1.
 CC KW Envelope protein; Hemagglutinin; Glycoprotein.
 CC FT NON TER 1 328
 CC FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
 CC FT CARBOHYD 7 550 HEMAGGLUTININ HA2 CHAIN.
 CC FT CARBOHYD 8 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CONFLICT 178 179 YV -> VI (IN PIR DATA BANK).
 CC FT CONFLICT 388 388
 CC SQ SEQUENCE 550 AA; 61856 MW; 48401C867A15BFC CRC64;
 CC
 CC Query Match 55.1%; Score 134; DB 1; Length 550;
 CC Best Local Similarity 100.0%; Pred. No. 2.2e-07;
 CC Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 CC
 CC 2 GLFGAIGFIENGWEGMIDWYG 24
 CC DB 330 GLFGAIGFIENGWEGMIDWYG 352
 CC
 CC RESULT 6
 CC HEMA_IADH6 STANDARD; PRT; 550 AA.
 CC ID HEMA_IADH6 Q84017;
 CC AC P12587; Q84017;
 CC DT 01-OCT-1989 (Rel. 12, Created)
 CC DT 01-APR-1990 (Rel. 14, Last sequence update)
 CC DT 16-OCT-2001 (Rel. 40, Last annotation update)
 CC DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
 CC DE Hemagglutinin HA2 chain] (Fragment).
 CC GN HA.
 CC OS Influenza A virus (strain A/Duck/Hokkaido/9/85).
 CC OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 CC CC Influenza A viruses; Influenzavirus A.
 CC OX NCBI_TaxID=11362;
 CC RN [1]
 CC RP SEQUENCE FROM N.A.
 CC RX MEDLINE=87265458; PubMed=2440178;
 CC RA Kida H., Kawaka Y., Nave C.W., Webster R.G.;
 CC RT "Antigenic and genetic conservation of H3 influenza virus in wild
 CC ducks.";
 CC RL Virology 159:109-119(1987).
 CC CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION.
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
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CC EMBL; M16742; AAA43148.1; -

DR PIR; P27813; HMI98.

DR HSSP; P03437; 1HGJ.

DR InterPro; IPR001364; Hemagglutn.

DR Pfam; PF00509; Hemagglutinin; 1.

DR PRINTS; PR00329; HEMAGGLUTN12.

DR Prodom; PD000225; Hemagglutn; 1.

KM Envelope protein; Hemagglutinin; Glycoprotein.

FT NON TER 1

FT CHAIN 1

FT CARBOHYD 330 550 HEMAGGLUTININ HAI CHAIN.

FT CARBOHYD 22 22 HEMAGGLUTININ HAZ CHAIN.

FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CONFLICT 8 8 Y -> N (IN PIR DATA BANK).

FT SEQUENCE 550 AA; 61711 MW; 67BCDB5F44736CFE CRC64;

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 2.2e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAIAGFIENGWEGMIDGWYG 24
|||
330 GLFGAIAGFIENGWEGMIDGWYG 352

Db

RESULT 7

HEMA_IADH7 STANDARD; PRT; 550 AA.

ID HEMA_IADH7

AC P12588; Q84018; Q89470;

DT 01-OCT-1989 (Rel. 12, Created)

DT 01-OCT-1989 (Rel. 12, Last sequence update)

DT 16-OCT-2001 (Rel. 40, Last annotation update)

DE Hemagglutinin precursor [Contains: Hemagglutinin HAI chain; Hemagglutinin HAZ chain] (Fragment).

GN HA.

OS Influenza A virus (strain A/Duck/Hokkaido/10/85).

OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;

CC Influenza A viruses; Influenzavirus A.

OX NCBI_TaxId=11363;

RN NCB1

RP SEQUENCE FROM N.A.

RA MEDLINE=87265458; PubMed=2440178;

RA Kida H., Kawacka Y., Naave C.W., Webster R.G.;

RT "Antigenic and genetic conservation of H3 influenza virus in wild ducks."

RT Viology 159:109-119(1987).

CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO CELL RECEPTORS AND FOR INITIATING INFECTION.

CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS (HA1 AND HA2) LINKED BY A DISULFIDE BOND.

CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.

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CC EMBL; M16743; AAA43149.1; -

DR HSSP; P03437; 3HMG.

DR InterPro; IPR001364; Hemagglutn.

DR Pfam; PF00509; Hemagglutinin; 1.

DR PRINTS; PR00329; HEMAGGLUTN12.

DR Prodom; PD000225; Hemagglutn; 1.

KM Envelope protein; Hemagglutinin; Glycoprotein.

FT NON TER 1

FT CHAIN 1

FT CARBOHYD 330 550 HEMAGGLUTININ HAI CHAIN.

FT CARBOHYD 22 22 HEMAGGLUTININ HAZ CHAIN.

FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT SEQUENCE 550 AA; 61761 MW; 68P81793281D53EB CRC64;

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 2.2e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAIAGFIENGWEGMIDGWYG 24
|||
330 GLFGAIAGFIENGWEGMIDGWYG 352

Db

RESULT 8

HEMA_IADHK STANDARD; PRT; 550 AA.

ID HEMA_IADHK

AC P43257;

DT 01-NOV-1995 (Rel. 32, Created)

DT 01-NOV-1995 (Rel. 32, Last sequence update)

DT 16-OCT-2001 (Rel. 40, Last annotation update)

DE Hemagglutinin precursor [Contains: Hemagglutinin HAI chain; Hemagglutinin HAZ chain] (Fragment).

GN HA.

OS Influenza A virus (strain A/Duck/Hong Kong/7/75).

OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;

CC Influenza A viruses; Influenzavirus A.

OX NCBI_TaxId=11364;

RN NCB1

RP SEQUENCE FROM N.A.

RA MEDLINE=91341491; PubMed=1875195;

RA Yasuda J., Shortridge K.F., Shimizu Y., Kida H.;

RT "Molecular evidence for a role of domestic ducks in the introduction of avian H3 influenza viruses to pigs in southern China, where the

RT of avian H3 influenza viruses to pigs in southern China, where the

RT J. Gen. Virol. 72:2007-2010(1991).

CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO CELL RECEPTORS AND FOR INITIATING INFECTION.

CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS (HA1 AND HA2) LINKED BY A DISULFIDE BOND.

CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.

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CC EMBL; D00929; BA00769.1; -

DR HSSP; P03437; 2V1U.

DR InterPro; IPR001364; Hemagglutn.

DR Pfam; PF00509; Hemagglutinin; 1.

DR PRINTS; PR00329; HEMAGGLUTN12.

DR Prodom; PD000225; Hemagglutn; 1.

KM Envelope protein; Hemagglutinin; Glycoprotein.

FT NON TER 1

FT CHAIN 1

FT CARBOHYD 330 550 HEMAGGLUTININ HAI CHAIN.

FT CARBOHYD 22 22 HEMAGGLUTININ HAZ CHAIN.

FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).

FT SEQUENCE 550 AA; 61549 MW; 864639B829F81BA9 CRC64;

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 2,2e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAFGIENGWEGMIDWYG 24
Db 330 GLFGAIAFGIENGWEGMIDWYG 352

RESULT 9

HEMA_IADHL STANDARD; PRT; 550 AA.
AC P43258;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
DE Hemagglutinin HA2 chain] (Fragment).
GN HA.
OS Influenza A virus (strain A/Duck/Hong Kong/64/76).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=45412;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=91341491; PubMed=1875195;
RA Yaeda J., Shortridge K.F., Shimizu Y., Kida H.;
RT "Molecular evidence for a role of domestic ducks in the introduction
RT of avian H3 influenza viruses to pigs in southern China, where the
RT A/Hong Kong/68 (H3N2) strain emerged.";
RT J. Gen. Virol. 72:2007-2010(1991).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
CC -----
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CC -----
CC EMBL; D00931; BAA00771.1; -.
CC HSSP; P03437; 2VU.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Hemagglutinin; Glycoprotein.
FT NON TER 1
FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CARBOHYD 8 550 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 22 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 550 AA; 61718 MW; A351C56789B4BE9A CRC64;

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 2,2e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAFGIENGWEGMIDWYG 24
Db 330 GLFGAIAFGIENGWEGMIDWYG 352

RESULT 10

HEMA_IAGHK STANDARD; PRT; 550 AA.
AC P43260;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
DE Hemagglutinin HA2 chain] (Fragment).
GN HA.
OS Influenza A virus (strain A/Goose/Hong Kong/10/76).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=45414;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=91341491; PubMed=1875195;
RA Yaeda J., Shortridge K.F., Shimizu Y., Kida H.;
RT "Molecular evidence for a role of domestic ducks in the introduction
RT of avian H3 influenza viruses to pigs in southern China, where the
RT A/Hong Kong/68 (H3N2) strain emerged.";
RT J. Gen. Virol. 72:2007-2010(1991).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; D00930; BAA00770.1; -.
CC HSSP; P03437; 2VU.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Hemagglutinin; Glycoprotein.
FT NON TER 1
FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
FT CARBOHYD 8 550 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 22 8 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 550 AA; 61676 MW; 9A1E094DA28BACD2 CRC64;

Query Match 55.1%; Score 134; DB 1; Length 550;
Best Local Similarity 100.0%; Pred. No. 2,2e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAFGIENGWEGMIDWYG 24
Db 330 GLFGAIAFGIENGWEGMIDWYG 352

RESULT 11

HEMA_IAGH2 STANDARD; PRT; 550 AA.
AC P11133; Q84019; Q84020;
DT 01-JUL-1989 (Rel. 11, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin [Contains: Hemagglutinin HA1 chain; Hemagglutinin HA2
DE chain] (Fragment).
GN HA.

OS Influenza A virus (strain A/Swine/Hong Kong/81/78).
 CC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 CC Influenza A viruses; Influenzavirus A.
 CC NCB1_Taxid=11497;
 CC [1]
 RN SEQUENCE FROM N.A.
 RP MEDLINE=88101364; Pubmed=3336940;
 RA Kida H., Shortridge K.F., Webster R.G.;
 RT "Origin of the hemagglutinin gene of H3N2 influenza viruses from pigs
 in China.";
 RT Virology 162:160-166(1988).
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION.
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
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 CC -----
 CC DR EMBL: M19057; AAA43212.1; -
 CC DR HSRP: P03437; 2V1U.
 CC DR InterPro: IPR001364; Hemagglutn.
 CC DR Pfam: PF00509; Hemagglutinin; 1.
 CC DR PRINTS: PR00329; HEMAGGLUTN12.
 CC DR ProDom: PD000225; Hemagglutn; 1.
 CC DR Hemagglutinin; Envelope protein; Glycoprotein.
 CC KW NON_TER 1 1
 CC FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
 CC FT CARBOHYD 8 550 HEMAGGLUTININ HA2 CHAIN.
 CC FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC SQ SEQUENCE 550 AA; 61437 MW; 1F2A7E758C531CE8 CRC64;
 CC
 CC Query Match 55.1%; Score 134; DB 1; Length 550;
 CC Best Local Similarity 100.0%; Pred. No. 2.2e-07;
 CC Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 CC
 CC QY 2 GLFGAIAGFIENGWGMIDGMYG 24
 CC DB 330 GLFGAIAGFIENGWGMIDGMYG 352
 CC
 CC RESULT 12
 CC HEMA_IASH3 STANDARD; PRT; 550 AA.
 CC ID P1134; Q84025; Q84026;
 CC AC 01-JUL-1988 (Rel. 11, Created)
 CC DT 01-JUL-1988 (Rel. 11, Last sequence update)
 CC DT 16-OCT-2001 (Rel. 40, Last annotation update)
 CC DE Hemagglutinin [Contains: Hemagglutinin HA1 chain; Hemagglutinin HA2
 CC chain] (Fragment).
 CC GN HA.
 CC OS Influenza A virus (strain A/Swine/Hong Kong/126/82).
 CC OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 CC CC Influenza A viruses; Influenzavirus A.
 CC NCBI_Taxid=11498;
 CC [1]
 RN SEQUENCE FROM N.A.
 RP MEDLINE=88101364; Pubmed=3336940;
 RA Kida H., Shortridge K.F., Webster R.G.;
 RT "Origin of the hemagglutinin gene of H3N2 influenza viruses from pigs
 RT in China.";
 RT Virology 162:160-166(1988).

CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION.
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
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 CC -----
 CC DR EMBL: M19056; AAA43211.1; ALT_TERM.
 CC DR HSRP: P03437; 2V1U.
 CC DR InterPro: IPR001364; Hemagglutn.
 CC DR Pfam: PF00509; Hemagglutinin; 1.
 CC DR PRINTS: PR00329; HEMAGGLUTN12.
 CC DR ProDom: PD000225; Hemagglutn; 1.
 CC DR Hemagglutinin; Envelope protein; Glycoprotein.
 CC KW NON_TER 1 1
 CC FT CHAIN 1 328 HEMAGGLUTININ HA1 CHAIN.
 CC FT CARBOHYD 8 550 HEMAGGLUTININ HA2 CHAIN.
 CC FT CARBOHYD 22 22 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 165 165 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 285 285 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC FT CARBOHYD 483 483 N-LINKED (GLCNAC. . .) (POTENTIAL).
 CC SQ SEQUENCE 550 AA; 61580 MW; 991F6D8BC02F24F2 CRC64;
 CC
 CC Query Match 55.1%; Score 134; DB 1; Length 550;
 CC Best Local Similarity 100.0%; Pred. No. 2.2e-07;
 CC Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 CC
 CC QY 2 GLFGAIAGFIENGWGMIDGMYG 24
 CC DB 330 GLFGAIAGFIENGWGMIDGMYG 352
 CC
 CC RESULT 13
 CC HEMA_IAMIC STANDARD; PRT; 566 AA.
 CC ID P03437;
 CC AC 21-JUL-1986 (Rel. 01, Created)
 CC DT 21-JUL-1986 (Rel. 01, Last sequence update)
 CC DT 15-SEP-2003 (Rel. 42, Last annotation update)
 CC DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
 CC Hemagglutinin HA2 chain].
 CC GN HA.
 CC OS Influenza A virus (strain A/Aichi/2/68).
 CC OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 CC CC Influenza A viruses; Influenzavirus A.
 CC NCBI_Taxid=150147;
 CC [1]
 RN SEQUENCE FROM N.A.
 RP MEDLINE=80254693; Pubmed=7402351;
 RA Verhoeven M., Fang R., Min Jou W., Devos R., Huylbrouck D.,
 RA Saman B., Fiers W.;
 RT "Antigenic drift between the haemagglutinin of the Hong Kong
 RT influenza strains A/Aichi/2/68 and A/Victoria/3/75.";
 RT Nature 286:771-776(1980).
 CC [2]
 RN X-RAY CRYSTALLOGRAPHY (3.0 ANGSTROMS).
 RP MEDLINE=81123029; Pubmed=7464906;
 RA Wilson I.A., Skehel J.D., Wiley D.C.;
 RT "Structure of the haemagglutinin membrane glycoprotein of influenza
 RT virus at 3-A resolution.";
 RT Nature 289:366-373(1981).
 CC [3]
 RN X-RAY CRYSTALLOGRAPHY.
 RP MEDLINE=88232903; Pubmed=3374584;

FT	TURN	350	350
FT	STRAND	351	351
FT	TURN	352	354
FT	STRAND	355	355
FT	STRAND	359	360
FT	TURN	361	361
FT	STRAND	367	373
FT	TURN	374	375
FT	STRAND	376	382
FT	HELIX	383	400
FT	TURN	401	401
FT	STRAND	406	407
FT	HELIX	421	471
FT	HELIX	472	474
FT	STRAND	475	477
FT	STRAND	482	485
FT	HELIX	491	498
FT	TURN	499	500
FT	HELIX	504	515
FT	TURN	518	519
SQ	SEQUENCE	566 AA;	63415 MW; E395659C23CAFECA CRC64;

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Query March 55.1%; Score 134; DB 1; Length 566;
Best Local Similarity 100.0%; Pred. NO. 2.3e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0.

Cy 2 GLFGAAGFIENGWEGMIDWYG 24
Db 346 GLFGAAGFIENGWEGMIDWYG 368

RESULT 14
HEMA_IADAJ
ID HEMA_IADAJ STANDARD; PRT; 566 AA.
AC P26134;
DT 01-MAY-1992 (Rel. 22, Created)
DT 01-MAY-1992 (Rel. 22, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Hemagglutinin precursor [Contains: Hemagglutinin HA1 chain;
  Hemagglutinin HA2 chain].
GN HA.
OS Influenza A virus (strain A/Duck/Alberta/76/76).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
ON NCBI_TaxID=113348;
RX [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=92114135; Pubmed=1731092;
RA Bean W.J., Schnell M., Katz J., Kanaoka Y., Naeye C., Gorman O.,
RA Webster R.G.;
RT "Evolution of the H3 influenza virus hemagglutinin from human and
  nonhuman hosts.";
RL J. Virol. 66:1129-1138(1992).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
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CC -----
DR EMBL; M73771; -; NOT_ANNOTATED_CDS.
DR HSSP; P03437; 2V1U.
DR InterPro; IPR001364; Hemagglu.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HemagglutN12.
DR ProDom; PD000225; Hemagglutn; 1.

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[illegible]

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QY      2  GLFGALGFIENGWEGMIDGWYG 24
      |||
Db      346  GLFGALGFIENGWEGMIDGWYG 368

Query March 15, 1997, Score 134; DB 1; Length 566;
Best Local Similarity 100.0%; Pred. No. 2,3e-07;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

RESULT 15
HEMA_IADU3 STANDARD; PRT; 566 AA.
AC P03442;
DT 21-JUL-1986 (Rel. 01, Last Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Hemagglutinin Precursor [Contains: Hemagglutinin HAI chain;
DE Hemagglutinin HA2 chain].
GN HA.
OS Influenza A virus (strain A/Duck/Ukraine/1/63).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCBI_TaxID=11374;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=82025542; PubMed=6169439;
RA Fang R., Min Jou W., Huybrecock D., Devos R., Fiers W.;
RT "Complete structure of A/Duck/Ukraine/63 influenza hemagglutinin
RT gene: animal virus as progenitor of human H3 Hong Kong 1968 influenza
RT hemagglutinin."
RL Cell 25:315-323 (1981).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION.
CC -1- SUBUNIT: HOMOTYMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND.
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
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CC -----
DR EMBL, J01087; CAA24271.1; -.
DR PDB, 1IBN; 08-AUG-01.
DR PDB, 1IBO; 08-AUG-01.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR ProDom; PD000225; Hemagglutn; 1.
KW Envelope protein; Hemagglutinin; Glycoprotein; Signal; 3D-structure.
FT SIGNAL 1 16
FT CHAIN 17 344 HEMAGGLUTININ HAI CHAIN.
FT CHAIN 346 566 HEMAGGLUTININ HA2 CHAIN.
FT CARBOHYD 24 24 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 38 38 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 54 54 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 97 97 N-LINKED (GLCNAC. . .) (POTENTIAL).

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FT CARBOHYD 181 181 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 301 301 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 499 499 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 566 AA; 63530 MW; E70F87F0AE1178F4 CRC64;

Query Match 55.1%; Score 134; DB 1; Length 566;
 Best Local Similarity 100.0%; Pred. No. 2.3e-07;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 GLFGAIAGFIEGMEGMDIGWYG 24
 |||||
 Db 346 GLFGAIAGFIEGMEGMDIGWYG 368

Search completed: January 30, 2004, 00:20:46
 Job time : 9.91549 secs

GenCore version 5.1.6
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OW protein - protein search, using sw model

Run on: January 29, 2004, 23:58:52 ; Search time 45.446 Seconds
(without alignments)
249.842 Million cell updates/sec

Title: US-09-461-684C-5
Perfect score: 243
Sequence: 1 CGLFGAIAGFIENGWEGMID.....KXXXXXXXXXXXXXXXXX 44

Scoring table: BLAST62
Gapop 10.0 , Gapept 0.5

Searched: 830525 seqs, 258052604 residues
Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

- 1: SP archaea:*
- 2: SP bacteria:*
- 3: SP fungi:*
- 4: SP human:*
- 5: SP_invertebrate:*
- 6: SP_mammal:*
- 7: SP_mmc:*
- 8: SP_organelle:*
- 9: SP_phage:*
- 10: SP_plant:*
- 11: SP_rodent:*
- 12: SP_virus:*
- 13: SP_vertebrate:*
- 14: SP_unclassified:*
- 15: SP_virus:*
- 16: SP_bacteriap:*
- 17: SP_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	134	55.1	384	12 Q8UK63	Q8UK63 Influenza a
2	134	55.1	566	12 Q98052	Q98052 Influenzavi
3	134	55.1	566	12 Q8U251	Q8U251 Influenza a
4	134	55.1	566	12 Q8OLN8	Q8OLN8 Influenza a
5	134	55.1	566	12 Q67132	Q67132 Influenza a
6	134	55.1	566	12 Q67135	Q67135 Influenzavi
7	134	55.1	566	12 Q8UXK3	Q8UXK3 Influenza a
8	134	55.1	566	12 Q91MA7	Q91MA7 Influenza a
9	134	55.1	566	12 Q9DHG0	Q9DHG0 Influenza a
10	134	55.1	566	12 Q910M5	Q910M5 Influenza a
11	134	55.1	566	12 Q67126	Q67126 Influenza a
12	133	54.7	301	12 Q9DXK3	Q9DXK3 Influenza a
13	132	54.3	550	12 Q82498	Q82498 Influenzavi
14	132	54.3	550	12 Q82498	Q82498 Influenzavi
15	132	54.3	550	12 Q82753	Q82753 Influenza v
16	132	54.3	566	12 Q82496	Q82496 Influenzavi

17	132	54.3	571	12 Q03909	Q03909 Influenza a
18	131	53.9	109	12 Q67050	Q67050 Influenzavi
19	131	53.9	109	12 Q67053	Q67053 Influenzavi
20	131	53.9	109	12 Q67051	Q67051 Influenzavi
21	131	53.9	109	12 Q67052	Q67052 Influenzavi
22	131	53.9	109	12 Q67053	Q67053 Influenzavi
23	131	53.9	109	12 Q67052	Q67052 Influenzavi
24	131	53.9	109	12 Q67053	Q67053 Influenzavi
25	131	53.9	109	12 Q67052	Q67052 Influenzavi
26	131	53.9	109	12 Q67053	Q67053 Influenzavi
27	131	53.9	109	12 Q67052	Q67052 Influenzavi
28	131	53.9	109	12 Q67053	Q67053 Influenzavi
29	131	53.9	109	12 Q67052	Q67052 Influenzavi
30	131	53.9	109	12 Q67053	Q67053 Influenzavi
31	131	53.9	109	12 Q67052	Q67052 Influenzavi
32	131	53.9	109	12 Q67053	Q67053 Influenzavi
33	131	53.9	109	12 Q67052	Q67052 Influenzavi
34	131	53.9	109	12 Q67053	Q67053 Influenzavi
35	131	53.9	109	12 Q67052	Q67052 Influenzavi
36	131	53.9	109	12 Q67053	Q67053 Influenzavi
37	131	53.9	109	12 Q67052	Q67052 Influenzavi
38	131	53.9	109	12 Q67053	Q67053 Influenzavi
39	131	53.9	109	12 Q67052	Q67052 Influenzavi
40	131	53.9	109	12 Q67053	Q67053 Influenzavi
41	131	53.9	109	12 Q67052	Q67052 Influenzavi
42	131	53.9	109	12 Q67053	Q67053 Influenzavi
43	131	53.9	109	12 Q67052	Q67052 Influenzavi
44	131	53.9	109	12 Q67053	Q67053 Influenzavi
45	131	53.9	109	12 Q67052	Q67052 Influenzavi

ALIGNMENTS

RESULT 1

ID Q8UK63 PRELIMINARY; PRT; 384 AA.

AC Q8UK63;

DT 01-OCT-2002 (TREMBLrel. 22. Created)

DT 01-OCT-2002 (TREMBLrel. 22. Last sequence update)

DT 01-MAR-2003 (TREMBLrel. 23. Last annotation update)

DE Hemagglutinin (Fragment).

GN H3HA.

OS Influenza A virus (A/teal/Germany/W201r/01).

OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;

OC Viruses; ssRNA negative-strand viruses; Influenzavirus A.

OX NCBI_TaxID=205472;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN-A/teal/Germany/W201r/01;

RA Werner O., Starick E., Mueller T., Muehle R.;

RT "Characterisation of avian influenza virus isolates from wild birds from Germany."

RL Submitted (AUG-2002) to the EMBL/GenBank/DBJ databases.

CC -!- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).

CC -!- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).

CC -!- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.

CC EMBL; AJ506781; CAD44999.1; -.

DR InterPro: IPR001364; Hemagglutn.

DR Pfam: PF00509; Hemagglutinin; 1.

DR PRINTS; PR00329; HEMAGGLUTIN12.

DR PRODOM; PD000225; Hemagglutn; 1.

KW Envelope protein; Glycoprotein; Hemagglutinin.

FT NON TER 384

FT SEQUENCE 384 AA; 42076 MW; 459731795CASCE38 CRC64;

Query Match 55.1%; Score 134; DB 12; Length 384;

Best Local Similarity 100.0%; Pred. No. 9.5e-08;

Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAGFIENGWEGMIDWYG 24

DB 346 GLFGAIAAGFIENGWEGMIDGWYG 368

RESULT 2

Q98052 PRELIMINARY; PRT; 566 AA.
 AC 098052; 01-FEB-1997 (TREMBlrel. 02, Created)
 DT 01-FEB-1997 (TREMBlrel. 02, Last sequence update)
 DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
 DE Hemagglutinin precursor (Fragment).
 OS Influenzavirus A.
 CC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 CC Influenza A viruses.
 OK NCBI_TaxID=197911;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=81053698; PubMed=6253883;
 RA Both G.W., Sleight M.J.;
 RT "Complete nucleotide sequence of the haemagglutinin gene from a human
 influenza virus of the Hong Kong subtype.";
 RL Nucleic Acids Res. 8:2561-2575(1980).
 RN [2]
 RP SEQUENCE OF 17-344 FROM N.A.
 RX MEDLINE=81194918; PubMed=6164798;
 RA Sleight M.J., Both G.W., Underwood P.A., Bender V.J.;
 RT "Antigenic drift in the hemagglutinin of the Hong Kong influenza
 subtype: Correlation of amino acid changes with alterations in viral
 antigenicity.";
 RL J. Virol. 37:845-853(1981).
 RN [3]
 RP SEQUENCE OF 17-566 FROM N.A.
 RX MEDLINE=82033276; PubMed=6169843;
 RA Both G.W., Sleight M.J.;
 RT "Conservation and variation in the hemagglutinins of Hong Kong subtype
 influenza viruses during antigenic drift.";
 RL J. Virol. 39:845-853(1981).
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL: J02135; AAA43189.1; -.
 DR HSRP; P03437; IHGE.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR Prodom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin; Signal.
 FT SIGNAL 1 16
 FT CHAIN 17 344
 FT CHAIN 346 566
 FT CHAIN 566 63414 MW; C447FDP65BE4FCF9 CRC64;
 SQ SEQUENCE 566 AA; 63414 MW; C447FDP65BE4FCF9 CRC64;
 Query Match 55.1%; Score 134; DB 12; Length 566;
 Best Local Similarity 100.0%; Pred. No. 1.4e-07;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 2 GLFGAIAAGFIENGWEGMIDGWYG 24
 DB 346 GLFGAIAAGFIENGWEGMIDGWYG 368
 RESULT 3
 Q8U251 PRELIMINARY; PRT; 566 AA.
 AC 08U251; 01-MAR-2002 (TREMBlrel. 20, Created)
 DT 01-MAR-2002 (TREMBlrel. 20, Last sequence update)
 DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
 DE Haemagglutinin.
 GN HA.

OS Influenza A virus (A/pet bird/Hong Kong/1559/99 (H3N8)).
 CC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 CC Influenza A viruses; Influenzavirus A.
 OK NCBI_TaxID=183665;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A/pet bird/Hong Kong/1559/99;
 RA Chin P., Shortridge K.F.;
 RT "Characterisation of avian H3 influenza viruses.";
 RL Submitted (JAN-2002) to the EMBL/Genbank/DBJ databases.
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; AJ427304; CAD20336.1; -.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR Prodom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 SQ SEQUENCE 566 AA; 63403 MW; F11C91B6A01B3484 CRC64;

Query Match 55.1%; Score 134; DB 12; Length 566;
 Best Local Similarity 100.0%; Pred. No. 1.4e-07;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAAGFIENGWEGMIDGWYG 24
 DB 346 GLFGAIAAGFIENGWEGMIDGWYG 368

RESULT 4

Q8QLN8 PRELIMINARY; PRT; 566 AA.
 AC Q8QLN8; 01-JUN-2002 (TREMBlrel. 21, Created)
 DT 01-JUN-2002 (TREMBlrel. 21, Last sequence update)
 DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
 DE Haemagglutinin.
 GN HA.
 OS Influenza A virus (A/aquatic bird/Hong Kong/399/99 (H3N8)).
 CC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 CC Influenza A viruses; Influenzavirus A.
 OK NCBI_TaxID=183664;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A/aquatic bird/Hong Kong/399/99;
 RA Chin P., Shortridge K.F.;
 RT "Characterisation of influenza viruses from wild aquatic birds.";
 RL Submitted (JAN-2002) to the EMBL/Genbank/DBJ databases.
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; AJ427297; CAD20332.1; -.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR Prodom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 SQ SEQUENCE 566 AA; 63412 MW; 68913C222C97B92E CRC64;

Query Match 55.1%; Score 134; DB 12; Length 566;
 Best Local Similarity 100.0%; Pred. No. 1.4e-07;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 GLFGAIAAGFIENGWEGMIDGWYG 24
 DB 346 GLFGAIAAGFIENGWEGMIDGWYG 368

RESULT 5

Q67132
ID 067132 PRELIMINARY; PRT; 566 AA.
AC 067132;
DT 01-NOV-1996 (Tremblrel. 01, Created)
DT 01-NOV-1996 (Tremblrel. 01, last sequence update)
DT 01-OCT-2002 (Tremblrel. 22, last annotation update)
DE Hemagglutinin.
GN HA.
OS Influenza A virus (strain A/Aichi/2/68).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCB1_TaxID=150147;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A/Aichi/2/68;
RA Min J.W., Verhoeven M., Fang R.-X., Devos R., Huybrecock D.,
RA Fiers W.,
RT "Shift and drift in influenza viruses."
RL (In) Carlile M.J., Collins J.F., Moseley B.E.B. (eds.);
RL SYMPOSIUM OF THE SOCIETY FOR GENERAL MICROBIOLOGY, pp.285-311,
RL Cambridge University Press, New York (1981)
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL; M55059; AAA43239.1; -.
DR HSSP; P03437; IHGE.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin.1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR Prodom; PD000225; Hemagglutn.1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
FT CHAIN 1 344 HEMAGGLUTININ.
FT CHAIN 2 566 NEURAMINIDASE.
SQ SEQUENCE 566 AA; 63441 MW; ESD1B97DP96FECA CRC64;
Query Match 55.1%; Score 134; DB 12; Length 566;
Best Local Similarity 100.0%; Pred. No. 1.4e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 GLFGAIGFTENGWEGMIDGMYG 24
Db 346 GLFGAIGFTENGWEGMIDGMYG 368
RESULT 6
Q67125
ID 067125 PRELIMINARY; PRT; 566 AA.
AC 067125;
DT 01-NOV-1996 (Tremblrel. 01, Created)
DT 01-NOV-1996 (Tremblrel. 01, last sequence update)
DT 01-OCT-2002 (Tremblrel. 22, last annotation update)
DE Hemagglutinin.
GN HA.
OS Influenzavirus A.
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses.
OX NCB1_TaxID=197911;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A/Seal/MA/3911/92;
RA MEDLINE=95146951; PubMed=7844533;
RA Callan R.J., Early G., Kida H., Hinshaw V.S.;
RT "The appearance of H3 influenza viruses in seals."
J. Gen. Virol. 76:199-203 (1995).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL; L31949; AAA64229.1; -.
DR HSSP; P03437; 2V1U.

DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin.1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR Prodom; PD000225; Hemagglutn.1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
SQ SEQUENCE 566 AA; 63456 MW; AE556302A9EBB99F CRC64;
Query Match 55.1%; Score 134; DB 12; Length 566;
Best Local Similarity 100.0%; Pred. No. 1.4e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 GLFGAIGFTENGWEGMIDGMYG 24
Db 346 GLFGAIGFTENGWEGMIDGMYG 368
RESULT 7
Q8UXR3
ID 08UXR3 PRELIMINARY; PRT; 566 AA.
AC 08UXR3;
DT 01-MAR-2002 (Tremblrel. 20, Created)
DT 01-MAR-2002 (Tremblrel. 20, last sequence update)
DT 01-OCT-2002 (Tremblrel. 22, last annotation update)
DE Hemagglutinin.
GN HA.
OS Influenza A virus (A/swine/Potsdam/35/82 (H3N2)).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCB1_TaxID=183769;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A/swine/Potsdam/35/82;
RA Groetinger I., Sues U., Groetinger C.;
RT "Evolution of european human and porcine influenza viruses."
Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
RL -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL; AJ252132; CAC81018.1; -.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin.1.
DR PRINTS; PR00329; HEMAGGLUTN12.
DR Prodom; PD000225; Hemagglutn.1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
SQ SEQUENCE 566 AA; 63529 MW; 6AA44C84B4DDE68A CRC64;
Query Match 55.1%; Score 134; DB 12; Length 566;
Best Local Similarity 100.0%; Pred. No. 1.4e-07;
Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 GLFGAIGFTENGWEGMIDGMYG 24
Db 346 GLFGAIGFTENGWEGMIDGMYG 368
RESULT 8
Q91MA7
ID 091MA7 PRELIMINARY; PRT; 566 AA.
AC 091MA7;
DT 01-DEC-2001 (Tremblrel. 19, Created)
DT 01-DEC-2001 (Tremblrel. 19, last sequence update)
DT 01-MAR-2003 (Tremblrel. 23, last annotation update)
DE Hemagglutinin.
OS Influenza A virus (A/Hong Kong/1/68 (H3N2)).
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza A viruses; Influenzavirus A.
OX NCB1_TaxID=108859;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A/Hong Kong/1/68 (H3N2);
MEDLINE=21287244; PubMed=11371620;

RA Brown E.G., Liu H., Kit L.C., Baird S., Neerallah M.;
 RT "Pattern of mutation in the genome of influenza A virus on adaptation
 RT to increased virulence in the mouse lung: identification of functional
 RT themes";
 RL Proc. Natl. Acad. Sci. U.S.A. 98:6883-6888(2001).
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; AF348176; AAKS1718.1; -.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR Prodom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 SQ SEQUENCE 566 AA; 63367 MW; 01BBD465BE158E1 CRC64;

Query Match 55.1%; Score 134; DB 12; Length 566;
 Best Local Similarity 100.0%; Pred. No. 1.4e-07;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAIGFIENGWEGMIDWYG 24
 DB 346 GLFGAIGFIENGWEGMIDWYG 368

RESULT 9
 ID Q9DHG0 PRELIMINARY; PRT; 566 AA.
 AC Q9DHG0;
 DT 01-MAR-2001 (TREMblrel. 16, Created)
 DT 01-MAR-2001 (TREMblrel. 16, Last sequence update)
 DT 01-OCT-2002 (TREMblrel. 22, Last annotation update)
 DE Haemagglutinin precursor.
 OS Influenza A virus H3N2.
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 CC Influenza A viruses; Influenzavirus A.
 CC NCBI_TaxID=41857;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=clone 7a;
 RA Mohsin M.A., Morris S.J., Smith H., Sweet C.;
 RT "Influenza virus-induced apoptosis: a dual role for viral
 RT neuraminidase";
 RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; AJ289703; CAC18525.1; -.
 DR HSP; P03437; 2V1U.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR Prodom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin; Signal.
 FT SIGNAL 16
 FT SIGNAL 16
 SQ SEQUENCE 566 AA; 63356 MW; 0BA681929300F72F CRC64;

Query Match 55.1%; Score 134; DB 12; Length 566;
 Best Local Similarity 100.0%; Pred. No. 1.4e-07;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAIGFIENGWEGMIDWYG 24
 DB 346 GLFGAIGFIENGWEGMIDWYG 368

RESULT 10
 ID Q910M5 PRELIMINARY; PRT; 566 AA.

AC Q910M5;
 DT 01-DEC-2001 (TREMblrel. 19, Created)
 DT 01-DEC-2001 (TREMblrel. 19, Last sequence update)
 DT 01-MAR-2003 (TREMblrel. 23, Last annotation update)
 DE Hemagglutinin.
 OS Influenza A virus (A/Hong Kong/1/68 (H3N2)).
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 CC Influenza A viruses; Influenzavirus A.
 CC NCBI_TaxID=108859;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A/Hong Kong/1/68 (H3N2);
 RX MEDLINE=21287244; PubMed=11371620;
 RA Brown E.G., Liu H., Kit L.C., Baird S., Neerallah M.;
 RT "Pattern of mutation in the genome of influenza A virus on adaptation
 RT to increased virulence in the mouse lung: identification of functional
 RT themes";
 RL Proc. Natl. Acad. Sci. U.S.A. 98:6883-6888(2001).
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; AF348179; AAKS1721.1; -.
 DR EMBL; AF348177; AAKS1719.1; -.
 DR EMBL; AF348178; AAKS1720.1; -.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR Prodom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 SQ SEQUENCE 566 AA; 63530 MW; 7CB9F5BAF1EE9F CRC64;

Query Match 55.1%; Score 134; DB 12; Length 566;
 Best Local Similarity 100.0%; Pred. No. 1.4e-07;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAIGFIENGWEGMIDWYG 24
 DB 346 GLFGAIGFIENGWEGMIDWYG 368

RESULT 11
 ID Q67126 PRELIMINARY; PRT; 566 AA.
 AC Q67126;
 DT 01-NOV-1996 (TREMblrel. 01, Created)
 DT 01-NOV-1996 (TREMblrel. 01, Last sequence update)
 DT 01-OCT-2002 (TREMblrel. 22, Last annotation update)
 DE Hemagglutinin.
 GN HA.
 OS Influenzavirus A.
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 CC Influenza A viruses.
 CC NCBI_TaxID=197911;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A/Seal/M/3984/92;
 RX MEDLINE=95146951; PubMed=7844533;
 RA Callan R.J., Early G., Kida H., Hinshaw V.S.;
 RT "The appearance of H3 influenza viruses in seals";
 RT J. Gen. Virol. 76:199-203(1995).
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; L32024; AAB64228.1; -.
 DR HSP; P03437; 2V1U.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR Prodom; PD000225; Hemagglutn; 1.

KM Envelope protein; Glycoprotein; Hemagglutinin.
 SQ SEQUENCE 566 AA; 63441 MW; 590576CB4CEB7D08 CRC64;
 Query Match 55.1%; Score 134; DB 12; Length 566;
 Best Local Similarity 100.0%; Pred. No. 1.4e-07;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAAGFIENGWEGMIDGWYG 24
 |||||
 DB 346 GLFGAAGFIENGWEGMIDGWYG 368

RESULT 12
 O9DXE3 PRELIMINARY; PRT; 301 AA.
 AC O9DXE3; 01-MAR-2001 (TREMBlrel. 16, Created)
 DT 01-MAR-2001 (TREMBlrel. 16, Last sequence update)
 DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
 DE Hemagglutinin (Fragment).
 GN HA1.
 OS Influenza A virus (A/Shorebird/Taiwan/31-4/99).
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 OC Influenza A viruses; Influenzavirus A.
 OX NCBI_TaxID=140665;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A/Shorebird/Taiwan/31-4/99;
 RA Lee M.S., Cheng P.C., Shieh J.H., Cheng M.C., Lee L.H., Shieh H.K.;
 RT "Identification and subtyping of avian influenza virus by reverse
 transcription-polymerase chain reaction.";
 RL Submitted (OCT-2000) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; AF111750; AAG3016.1; -.
 DR InterPro; IPR001364; Hemagglutn;
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR ProDom; PD000225; HEMAGGLUTN; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 FT NON_TER 1
 FT NON_TER 301
 SQ SEQUENCE 301 AA; 32701 MW; 62A403758B764D57 CRC64;
 Query Match 54.7%; Score 133; DB 12; Length 301;
 Best Local Similarity 95.7%; Pred. No. 9.7e-08;
 Matches 22; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAAGFIENGWEGMIDGWYG 24
 |||||
 DB 250 GLFGAAGFIENGWEGMIDGWYG 272

RESULT 13
 O82499 PRELIMINARY; PRT; 550 AA.
 AC O82499;
 DT 01-NOV-1996 (TREMBlrel. 01, Created)
 DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
 DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
 DE Hemagglutinin HA1 and HA2 (Fragment).
 OS Influenzavirus A.
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 OC Influenza A viruses.
 OX NCBI_TaxID=197911;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A/Philippines/2/82/BS;
 RA Hartley C.A., Ward A.C., Anders B.M.;
 RT "Virulence of influenza virus for mice is associated with loss of

RT oligosaccharide from the hemagglutinin molecule.";
 RL Submitted (APR-1994) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; U08859; AAA18782.1; -.
 DR HSSP; P03437; 2YIU.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR ProDom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 FT NON_TER 1
 FT CHAIN 1
 FT CHAIN 328
 FT CHAIN 550
 SQ SEQUENCE 550 AA; 61772 MW; 50BD62B6BF11PD8 CRC64;
 Query Match 54.3%; Score 132; DB 12; Length 550;
 Best Local Similarity 95.7%; Pred. No. 2.3e-07;
 Matches 22; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 GLFGAAGFIENGWEGMIDGWYG 24
 |||||
 DB 330 GLFGAAGFIENGWEGMIDGWYG 352

RESULT 14
 O82498 PRELIMINARY; PRT; 550 AA.
 AC O82498;
 DT 01-NOV-1996 (TREMBlrel. 01, Created)
 DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
 DT 01-OCT-2002 (TREMBlrel. 22, Last annotation update)
 DE Hemagglutinin HA1 and HA2 (Fragment).
 OS Influenzavirus A.
 OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
 OC Influenza A viruses.
 OX NCBI_TaxID=197911;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=A/Philippines/2/82;
 RA Hartley C.A., Ward A.C., Anders B.M.;
 RT "Virulence of influenza virus for mice is associated with loss of
 oligosaccharide from the hemagglutinin molecule.";
 RL Submitted (APR-1994) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
 CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
 CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
 (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
 CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
 DR EMBL; U08858; AAA18781.1; -.
 DR HSSP; P03437; 2YIU.
 DR InterPro; IPR001364; Hemagglutn.
 DR Pfam; PF00509; Hemagglutinin; 1.
 DR PRINTS; PR00329; HEMAGGLUTN12.
 DR ProDom; PD000225; Hemagglutn; 1.
 KW Envelope protein; Glycoprotein; Hemagglutinin.
 FT NON_TER 1
 FT CHAIN 1
 FT CHAIN 328
 FT CHAIN 550
 SQ SEQUENCE 550 AA; 61802 MW; 114413B1CE5A1F6A CRC64;
 Query Match 54.3%; Score 132; DB 12; Length 550;

Best Local Similarity 95.7%; Pred. No. 2.3e-07;
Matches 22; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 GFGAIAGFIENGEGMIDGWYG 24
|:|||||
Db 330 GFGAIAGFIENGEGMIDGWYG 352

RESULT 15

Q82753 PRELIMINARY; PRT; 550 AA.
AC Q82753;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
DE Haemagglutinin (Fragment).
OS Influenza virus.
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Unclassified Orthomyxoviridae.
OX NCBI_TaxID=11309;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A/Philippines/2/82/BS/ML10;
RX MEDLINE=97300854; PubMed=9155874;
RA Hartley C.A., Reading P.C., Ward A.C., Anders E.M.;
RT "Changes in the hemagglutinin molecule of influenza type A (H3N2)
RT virus associated with increased virulence for mice.";
RL Arch. Virol. 142:75-88 (1997).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=A/Philippines/2/82/BS/ML10;
RX MEDLINE=97456249; PubMed=9311563;
RA Ward A.C.;
RT "Virulence of Influenza A virus for mouse lung.";
RL Virus Genes 14:187-194 (1997).
CC -1- FUNCTION: HEMAGGLUTININ IS RESPONSIBLE FOR ATTACHING THE VIRUS TO
CC CELL RECEPTORS AND FOR INITIATING INFECTION (BY SIMILARITY).
CC -1- SUBUNIT: HOMOTRIMER. EACH OF THE MONOMERS IS FORMED BY TWO CHAINS
CC (HA1 AND HA2) LINKED BY A DISULFIDE BOND (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE INFLUENZA HEMAGGLUTININ FAMILY.
DR EMBL; U08905; AAC79579.1; -.
DR HSSP; P03437; 2VIT.
DR InterPro; IPR001364; Hemagglutn.
DR Pfam; PF00509; Hemagglutinin; 1.
DR PRINTS; PR00329; HEMAGGLUTIN12.
DR Prodom; PD000225; Hemagglutn; 1.
KW Envelope protein; Glycoprotein; Hemagglutinin.
FT NON TER 1
FT CHAIN 1 328 HAEMAGGLUTININ HA1.
FT CHAIN 330 550 HAEMAGGLUTININ HA2.
SQ SEQUENCE 550 AA; 61745 MW; 692A49DE678AC4BC CRC64;

Query Match 54.3%; Score 132; DB 12; Length 550;
Best Local Similarity 95.7%; Pred. No. 2.3e-07;
Matches 22; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 GFGAIAGFIENGEGMIDGWYG 24
|:|||||
Db 330 GFGAIAGFIENGEGMIDGWYG 352

Search completed: January 30, 2004, 00:24:41
Job time : 45.446 secs

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OM protein - protein search, using sw model

Run on: January 30, 2004, 07:06:28 ; Search time 21 Seconds
(without alignments)
36.636 Million cell updates/sec

Title: SEQ10
Perfect score: 38
Sequence: 1 sinifek1 8

Scoring table: BIOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283308 seqs, 96168682 residues

Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :
1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	38	100.0	386	1	OACH
2	34	89.5	627	2	S37954
3	33	86.8	168	2	P90095
4	33	86.8	315	2	T10818
5	33	86.8	383	2	S11433
6	32	84.2	260	2	G87349
7	32	84.2	347	2	A99989
8	32	84.2	1305	2	T23314
9	31	81.6	244	2	T06961
10	31	81.6	304	2	C64109
11	31	81.6	307	2	E71206
12	31	81.6	309	2	H90548
13	31	81.6	364	2	A84726
14	31	81.6	397	2	T34441
15	31	81.6	680	2	T42923
16	30	78.9	142	2	A83093
17	30	78.9	232	1	DXCH
18	30	78.9	347	2	A82358
19	30	78.9	349	2	H71923
20	30	78.9	378	2	T18486
21	30	78.9	440	2	F70117
22	30	78.9	467	2	AF1317
23	30	78.9	470	2	D84614
24	30	78.9	481	2	B70179
25	30	78.9	487	2	C97144
26	30	78.9	520	2	B82206
27	30	78.9	558	2	S61604
28	30	78.9	610	2	T25262
29	30	78.9	842	2	E72373

30	30	78.9	845	2	D90130	hypothetical prote
31	30	78.9	1004	2	B69483	hypothetical prote
32	30	78.9	1115	2	B84476	probable TPR repea
33	30	78.9	1163	2	G97236	ATPase involved in
34	30	78.9	2116	2	T49818	glutamate synthase
35	30	78.9	2469	2	H36812	hypothetical prote
36	30	78.9	4092	1	S38128	dynein heavy chain
37	29	76.3	69	2	A96966	protein PIN21.2 (1
38	29	76.3	88	4	S54022	hypothetical prote
39	29	76.3	110	2	F97094	probable transcrip
40	29	76.3	141	2	H69044	peptidylprolyl iso
41	29	76.3	143	2	S45537	peptidylprolyl iso
42	29	76.3	170	2	G68356	hypothetical prote
43	29	76.3	209	2	D86882	hypothetical prote
44	29	76.3	237	2	F64506	2-hydroxyhepta-2,4
45	29	76.3	275	1	C69030	M11225 protein hom

ALIGNMENTS

RESULT 1

OACH
Ovalbumin [validated] - chicken
C:Species: Gallus gallus (chicken)
C:Date: 31-Dec-1979 #sequence revision 30-Jun-1993 #text change 15-Sep-2000
C:Accession: A90455; I50402; I50605; A93197; A93827; A90092; A61297; A42793; A01
R:Woo, S.L.C.; Beattie, W.G.; Catterall, J.F.; Dugalczyk, A.; Staden, R.; Brownlee, G.G
Biochemistry 20, 6437-6446, 1981
A>Title: Complete nucleotide sequence of the chicken chromosomal ovalbumin gene and its
A:Reference number: A90455; MUID:82069038; PMID:6272839
A:Accession: A90455
A:Molecule type: DNA
A:Residues: 1-386 <WOO>
A:Cross-References: EMBL:V00438; NID:963719; PIDN:CAA23716.1; PID:G808974
A>Note: A number of silent polymorphic sites are identified and discussed
A:Note: Thr-188 is also predicted
R:Catterall, J.F.; O'Malley, B.W.; Robertson, M.A.; Staden, R.; Tanaka, Y.; Brownlee, G
Nature 275, 510-513, 1978
A>Title: Nucleotide sequence homology at 12 intron-exon junctions in the chick ovalbumin
A:Reference number: I50402; MUID:79010682; PMID:692731
A:Accession: I50402
A:Molecule type: DNA
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-386 <CAT>
A:Cross-References: GB:M34352; NID:G212501; PIDN:AAA48998.1; PID:G212503
R:Robertson, M.A.; Staden, R.; Tanaka, Y.; Catterall, J.F.; Brownlee, G.
Nature 278, 370-372, 1979
A>Title: Sequence of three introns in the chick ovalbumin gene.
A:Reference number: I50605; MUID:79135070; PMID:423993
A:Accession: I50605
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-4, 'A', '6-118, 'F', '120-155 <ROB>
A:Cross-References: EMBL:V00382; NID:963051; PIDN:CAA23681.1; PID:G63052
R:McReynolds, L.; O'Malley, B.W.; Nisbet, A.D.; Fothergill, J.E.; Givoli, D.; Fields, S.,
Nature 273, 723-728, 1978
A>Title: Sequence of chicken ovalbumin mRNA.
A:Reference number: A93197; MUID:78199842; PMID:661981
A:Accession: A93197
A:Molecule type: mRNA
A:Residues: 1-386 <MCB>
A:Cross-References: EMBL:V00383; NID:963053
A>Note: a minor component has Asp-312
R:Palmiter, R.D.; Gagnon, J.; Walsby, K.A.
Proc. Natl. Acad. Sci. U.S.A. 75, 94-98, 1978
A>Title: Ovalbumin: a secreted protein without a transient hydrophobic leader sequence.
A:Reference number: A93827; MUID:78116057; PMID:272676
A:Accession: A93827
A:Molecule type: protein
A:Residues: 2-33, 'X', '35-36 <PAL>
R:Thompson, E.O.P.; Fisher, W.K.
Aust. J. Biol. Sci. 31, 443-446, 1978

A>Title: A correction and extension of the acetylated amino terminal sequence of ovalbumin
 A:Reference number: A90093; PMID:79186958; PMID:751625
 A:Accession: A90093
 A:Molecule type: protein
 A:Residues: 2-17 <TH1>
 R:Thompson, E.O.P.; Fisher, W.K.
 Aust. J. Biol. Sci. 31, 433-442, 1978
 A>Title: Amino acid sequences containing half-cysteine residues in ovalbumin.
 A:Reference number: A90092; PMID:79186957; PMID:751624
 A:Accession: A90092
 A:Molecule type: protein
 A:Residues: 6-17,30-36,61-79,116-124,367-374,380-386 <TH2>
 R:Tsunawasa, S.; Narita, K.
 J. Biochem. 92, 607-613, 1982
 A>Title: Micro-identification of amino-terminal acetylaminic acids in proteins.
 A:Reference number: A61297; PMID:83056735; PMID:6554709
 A:Accession: A61297
 A:Molecule type: protein
 A:Residues: 2-6 <TSU>
 R:Takahashi, N.; Hirose, M.
 J. Biol. Chem. 267, 11565-11572, 1992
 A>Title: Reversible denaturation of disulfide-reduced ovalbumin and its reoxidation gene
 A:Reference number: A42793; PMID:92283876; PMID:1597484
 A:Accession: A42793
 A:Molecule type: protein
 A:Residues: 60-73, 'X', 75-85,112-119, 'EX',122-123 <TKA>
 R:Stein, P.E.; Leslie, A.G.W.
 submitted to the Brookhaven Protein Data Bank, November 1990
 A:Reference number: A50294; PDB:1OVA
 A:Contents: annotation; X-ray crystallography, 1.95 angstroms, residues 2-386
 R:Stein, P.E.; Leslie, A.G.W.; Finch, J.T.; Carrell, R.W.
 J. Mol. Biol. 221, 941-959, 1991
 A>Title: Crystal structure of unbleached ovalbumin at 1.95 Angstroms resolution.
 A:Reference number: A58761; PMID:92046044; PMID:1942038
 A:Contents: annotation; X-ray crystallography, 1.95 angstroms
 A:Genetics:
 A:Introns: 56/3; 73/3; 116/3; 156/1; 203/3; 255/3
 C:Superfamily: antithrombin III
 C:Keywords: acetylated amino end; glycoprotein; phosphoprotein
 F:2-384/Product: ovalbumin #status experimental <MAT>
 F:2/Modified site: acetylated amino end (Gly) (in mature form) #status experimental
 F:69,345/Binding site: phosphate (Ser) (covalent) #status experimental
 F:74-121/Disulfide bonds: #status experimental
 F:293/Binding site: carbohydrate (Asn) (covalent) #status experimental

Query Match 100.0%; Score 38; DB 1; Length 386;
 Best Local Similarity 100.0%; Pred. No. 2.3;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEXL 8
 DB 258 SIINFEXL 265

RESULT 2
 S37954
 RNA polymerase I transcription factor RRM3 - yeast (Saccharomyces cerevisiae)
 N:Alternate names: protein YKL125W
 C:Species: Saccharomyces cerevisiae
 C:Date: 03-May-1994 #sequence_revision 03-May-1994 #text_change 13-Mar-1998
 C:Accession: S37954; PMID:751600
 R:Ramazani Rad, M.; Xu, G.; Kirchrath, L.; Fritze, C.; Keuchel, H.; Hollenberg, C.P.
 submitted to the Protein Sequence Database, March 1994
 A:Reference number: S37953
 A:Accession: S37954
 A:Molecule type: DNA
 A:Residues: 1-627 <RML>
 A:Cross-references: EMBL:Z28125; NID:9486210; PID:9486211; MIPS:YKL125W
 A:Experimental source: strain S288C
 R:Yamamoto, R.T.; Nogi, Y.; Dodd, J.A.; Nomura, M.
 EMBO J. 15, 3964-3973, 1996
 A>Title: RRM3 gene of Saccharomyces cerevisiae encodes an essential RNA polymerase I tra
 A:Reference number: S71600; PMID:96324404; PMID:8670901

A:Accession: S71600
 A:Molecule type: DNA
 A:Residues: 1-627 <YAM>
 C:Genetics:
 A:Gene: SGD:RRM3
 A:Cross-references: SGD:S0001608; MIPS:YKL125W
 A:Map position: 11L
 C:Keywords: nucleus

Query Match 89.5%; Score 34; DB 2; Length 627;
 Best Local Similarity 100.0%; Pred. No. 28;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEXL 7
 DB 118 SIINFEXL 124

RESULT 3
 P90095
 hypothetical protein orf168 [imported] - Guillardia theta nucleomorph
 C:Species: nucleomorph Guillardia theta
 A>Note: a nucleomorph is the vestigial nucleus of a eukaryotic endosymbiont
 C:Date: 10-May-2001 #sequence_revision 10-May-2001 #text_change 24-May-2001
 C:Accession: P90095
 R:Douglas, S.; Zauner, S.; Fraunholz, M.; Beaton, M.; Penny, S.; Deng, L.T.; Wu, X.; Re
 Nature 410, 1091-1096, 2001
 A>Title: The highly reduced genome of an enslaved algal nucleus.
 A:Reference number: A99082; PMID:11323671; PMID:11323671
 A:Accession: P90095
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-168 <DOU>
 A:Cross-references: GB:AF165818; NID:913794518; PIDN:AAK39893.1; GSPDB:GN00150
 C:Genetics:
 A:Gene: orf168
 A:Map position: 1
 A:Genome: nucleomorph
 C:Keywords: nucleomorph

Query Match 86.8%; Score 33; DB 2; Length 168;
 Best Local Similarity 75.0%; Pred. No. 11;
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEXL 8
 DB 108 SIINFEXL 115

RESULT 4
 T10818
 1-aminocyclopropane-1-carboxylate oxidase (EC 1.4.3.-) - kidney bean
 C:Species: Phaseolus vulgaris (kidney bean)
 C:Date: 16-Jul-1999 #sequence_revision 16-Jul-1999 #text_change 19-May-2000
 C:Accession: T10818
 R:Pidgen, C.M.; Facchini, P.J.; Reid, D.M.
 submitted to the EMBL Data Library, March 1998
 A:Reference number: Z17172
 A:Accession: T10818
 A:Status: translated from GB/EMBL/DBJ
 A:Molecule type: mRNA
 A:Residues: 1-315 <PID>
 A:Cross-references: EMBL:AF053354; NID:93037046; PIDN:AAK12934.1; PID:93037047
 A:Experimental source: cultivar Taylor bush bean, leaf
 C:Genetics:
 A:Gene: ACO1
 C:Superfamily: 1-aminocyclopropane-1-carboxylate oxidase
 C:Keywords: ethylene biosynthesis; iron; metalloprotein; oxidoreductase
 F:39,177,23/Binding site: iron (His) #status predicted

Query Match 86.8%; Score 33; DB 2; Length 315;
 Best Local Similarity 85.7%; Pred. No. 22;
 Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 IINFEKL 8
:|||||
Db 6 VINFEKL 12

RESULT 5

ovalbumin - Japanese quail
S11433
C:Species: Coturnix coturnix japonica (Japanese quail)
C>Date: 21-Nov-1993 #sequence_revision 10-Nov-1995 #text_change 16-Jul-1999
C:Accession: S11433
R:Mucha, J.; Klaudiny, J.; Klaudinyova, V.; Hanes, J.; Simuth, J.
Nucleic Acids Res. 18, 5553, 1990
A>Title: The sequence of Japanese quail ovalbumin cDNA.
A:Reference number: S11433; MUID:91016850; PMID:2216734
A:Accession: S11433
A>Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-383 <MUC>
A:Cross-references: EMBL:X53964; NID:g62643; PIDN:CAA37916.1; PID:g62644
C:Superfamily: antithrombin III

Query Match 86.8%; Score 33; DB 2; Length 383;
Best Local Similarity 87.5%; Pred. No. 27;
Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEKL 8
:|||||
Db 258 STINFEKL 265

RESULT 6

conserved hypothetical protein CC0810 [imported] - Caulobacter crescentus
G87349
C:Species: Caulobacter crescentus
C>Date: 20-Apr-2001 #sequence_revision 20-Apr-2001 #text_change 20-Apr-2001
C:Accession: G87349
R:Niemann, W.C.; Feldblyum, T.V.; Paulsen, I.T.; Nelson, K.E.; Eisen, J.; Heidelberg, J.
B.; Laub, M.T.; Deboy, R.T.; Dodson, R.J.; Durkin, A.S.; Gwinn, M.L.; Haft, D.H.; Kojouhar, M.; Esmolaeva, M.; White, O.; Salzberg, S.L.; Shapiro, L.; Venter, J.C.; Fraser, C.M.
Proc. Natl. Acad. Sci. U.S.A. 98, 4136-4141, 2001
A>Title: Complete Genome Sequence of Caulobacter crescentus.
A:Reference number: A87249; MUID:21173698; PMID:11259647
A:Accession: G87349
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-260 <STO>
A:Cross-references: GB:AE005673; NID:g13422057; PIDN:AAK22795.1; GSPDB:GN00148
C:Genetics:

Query Match 84.2%; Score 32; DB 2; Length 260;
Best Local Similarity 87.5%; Pred. No. 29;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 STINFEKL 8
:|||||
Db 215 STINFEKL 222

RESULT 7

A99989
Cyclin B [imported] - Guillardia theta nucleomorph
C:Species: nucleomorph guillardia theta
A>Note: a nucleomorph is the vestigial nucleus of a eukaryotic endosymbiont
C>Date: 10-May-2001 #sequence_revision 10-May-2001 #text_change 24-May-2001
C:Accession: A99989
R:Douglas, S.; Zauner, S.; Fraunholz, M.; Beaton, M.; Penny, S.; Deng, L.T.; Wu, X.; Reif
Nature 410, 1091-1096, 2001
A>Title: The highly reduced genome of an enslaved algal nucleus.
A:Reference number: A99082; MUID:11323671; PMID:11323671
A:Accession: A99989

A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-347 <DOU>
A:Cross-references: GB:AF165818; NID:g13794469; PIDN:AAK39844.1; GSPDB:GN00150
C:Genetics:
A:Gene: cycB
A:Map position: 1
A:Genome: nucleomorph
C:Keywords: nucleomorph

Query Match 84.2%; Score 32; DB 2; Length 347;
Best Local Similarity 62.5%; Pred. No. 40;
Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEKL 8
:|||||
Db 86 NVINFEKL 93

RESULT 8

hypothetical protein T14G10.2 - Caenorhabditis elegans
T23314
C:Species: Caenorhabditis elegans
C>Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 29-Oct-1999
C:Accession: T23314; T24919
R:Wild, A.
submitted to the EMBL Data Library, February 1996
A:Reference number: Z19725
A:Accession: T23314
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-1305 <WIL>
A:Cross-references: EMBL:Z69664; PIDN:CAA3519.1; GSPDB:GN00022; CESP:T14G10.2
A:Experimental source: clone K04D7
R:Wild, A.
submitted to the EMBL Data Library, January 1996
A:Reference number: Z19954
A:Accession: T24919
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-1305 <WIL>
A:Cross-references: EMBL:Z68880; PIDN:CAA33100.1; GSPDB:GN00022; CESP:T14G10.2
A:Experimental source: clone T14G10
C:Genetics:
A:Gene: CESP:T14G10.2
A:Map position: 4
A:Introns: 450/1; 463/2; 696/2; 763/2; 843/2; 935/3; 1012/1; 1091/1; 1143/1; 1189/2; 125

Query Match 84.2%; Score 32; DB 2; Length 1305;
Best Local Similarity 85.7%; Pred. No. 1.7e+02;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 IINFEKL 8
:|||||
Db 903 IINFEKL 909

RESULT 9

T06961
ABC transport protein homolog - Cyanophora paradoxa cyanelle
C:Species: cyanelle Cyanophora paradoxa
C>Date: 30-Apr-1999 #sequence_revision 30-Apr-1999 #text_change 17-Mar-2000
C:Accession: T06961
R:Stewart, V.L.; Michalowski, C.B.; Luffelhardt, W.; Bohner, H.J.; Bryant, D.A.
submitted to the EMBL Data Library, July 1995
A>Description: Nucleotide sequence of the cyanelle genome from Cyanophora paradoxa.
A:Reference number: Z15840
A:Accession: T06961
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-244 <STI>
A:Cross-references: EMBL:U30821; NID:g1016083; PIDN:AAA81304.1; PID:g1016217
A:Experimental source: strain Pringsheim LB555

C:Genetics:
 A:Genome: cyanelle
 C:Superfamily: unassigned ATP-binding cassette proteins; ATP-binding cassette homology
 C:Keywords: cyanelle

Query Match 81.6%; Score 31; DB 2; Length 244;
 Best Local Similarity 75.0%; Pred. No. 44;
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEK 8
 :|||||:
 Db 204 STINFEK 211

RESULT 10
 C64109
 site-specific DNA-methyltransferase (cytosine-specific) (EC 2.1.1.73) - Haemophilus inf
 C:Species: Haemophilus influenzae
 C:Date: 18-Aug-1995 #sequence_revision 18-Aug-1995 #text_change 21-Jul-2000
 C:Accession: C64109
 R:Flieschmann, R.D.; Adams, M.D.; White, O.; Clayton, R.A.; Kirkness, E.F.; Kerlavage, A
 ; Gocayne, J.D.; Scott, J.; Shirley, R.; Liu, L.I.; Glodek, A.; Kelley, J.M.; Weidman, D
 ; D.M.; Brandon, R.C.; Fine, L.D.; Fritchman, J.L.; Fuhmann, J.L.; Geoghegan, N.S.M.
 Science 269, 496-512, 1995
 A:Authors: Gnehm, C.L.; McDonald, L.A.; Small, K.V.; Fraser, C.M.; Smith, H.O.; Venter,
 A:Title: Whole-genome random sequencing and assembly of Haemophilus influenzae Rd.
 A:Reference number: A64000; PMID:95350630; PMID:7542800
 A:Accession: C64109
 A:Status: nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-304 <TIGR>
 A:Cross-references: GB:142023; NID:G3212210; PIDN:AAC22700.1; PID:G1574073; T
 C:Superfamily: site-specific methyltransferase (cytosine-specific) EcoRII
 C:Keywords: methyltransferase; restriction modification system; S-adenosylmethionine

Query Match 81.6%; Score 31; DB 2; Length 304;
 Best Local Similarity 85.7%; Pred. No. 56;
 Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEK 7
 :|||||:
 Db 291 ATINFEK 297

RESULT 11
 E71206
 hypothetical protein PH1919 - Pyrococcus horikoshii
 C:Species: Pyrococcus horikoshii
 C:Date: 14-Aug-1998 #sequence_revision 14-Aug-1998 #text_change 21-Jul-2000
 C:Accession: E71206
 R:Kawarabayashi, Y.; Sawada, M.; Horikawa, H.; Hatakeyama, Y.; Hino, Y.; Yamamoto, S.; Seki
 N.; Ohfuku, Y.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; Kushida, N.; Oguchi
 DNA Res. 5, 55-76, 1998
 A:Title: Complete sequence and gene organization of the genome of a hyper-thermophilic a
 A:Reference number: A71000; PMID:98344137; PMID:9679194
 A:Accession: E71206
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-307 <RAW>
 A:Cross-references: GB:AP000007; NID:G3236134; PIDN:BA31044.1; PID:G3258361
 A:Experimental source: strain OT3
 A:Note: this accession replaces an interim accession for a sequence replaced by GenBank
 C:Genetics:
 A:Gene: PH1919

Query Match 81.6%; Score 31; DB 2; Length 307;
 Best Local Similarity 75.0%; Pred. No. 57;
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEK 8
 :|||||:
 Db 3 NINFEK 10

RESULT 12
 H90548
 hypothetical protein MYP2960 [imported] - Mycoplasma pulmonis (strain UAB CTIP)

C:Species: Mycoplasma pulmonis
 C:Date: 24-May-2001 #sequence_revision 24-May-2001 #text_change 03-Aug-2001
 C:Accession: H90548
 R:Chabaud, I.; Heilig, R.; Ferris, S.; Barbe, V.; Samson, D.; Galisson, F.; Moszer, I.;
 Nucleic Acids Res. 29, 2145-2153, 2001
 A:Title: The complete genome sequence of the murine respiratory pathogen Mycoplasma pulm
 A:Reference number: A99512; PMID:21267165; PMID:11353084
 A:Accession: H90548
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-309 <KUR>
 A:Cross-references: GB:AL445566; PID:G14089710; PIDN:CA13469.1; GSPDB:GN00153
 A:Experimental source: strain UAB CTIP
 C:Genetics:
 A:Gene: MYP2960
 A:Genetic code: SGC3

Query Match 81.6%; Score 31; DB 2; Length 309;
 Best Local Similarity 71.4%; Pred. No. 57;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEK 7
 :|||||:
 Db 287 STINFEK 293

RESULT 13
 A64726
 probable poly(ADP-ribose) glycohydrolase [imported] - Arabidopsis thaliana
 C:Species: Arabidopsis thaliana (mouse-ear cress)
 C:Date: 02-Feb-2001 #sequence_revision 02-Feb-2001 #text_change 02-Feb-2001
 C:Accession: A64726
 R:Lin, X.; Kaul, S.; Rounsley, S.D.; Shea, T.P.; Benito, M.I.; Town, C.D.; Fujii, C.Y.;
 M.; Koo, H.; Moffat, K.S.; Cronin, L.A.; Shen, M.; Vanaken, S.E.; Umayam, L.; Tallon, L.
 eus, D.; Nierman, W.C.; White, O.; Eilen, J.A.; Salzberg, S.L.; Fraser, C.M.; Venter, J
 Nature 402, 761-768, 1999
 A:Title: Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana.
 A:Reference number: A64420; PMID:20083487; PMID:10617197
 A:Accession: A64726
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-364 <STO>
 A:Cross-references: GB:AE002093; NID:G4887750; PIDN:AA032286.1; GSPDB:GN00139
 C:Genetics:
 A:Gene: At2g31860
 A:Map position: 2

Query Match 81.6%; Score 31; DB 2; Length 364;
 Best Local Similarity 85.7%; Pred. No. 68;
 Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 IINFEK 8
 :|||||:
 Db 88 IINFEK 94

RESULT 14
 T34441
 hypothetical protein K11H12.3 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans
 C:Date: 29-Oct-1999 #sequence_revision 29-Oct-1999 #text_change 29-Oct-1999
 C:Accession: T34441
 R:Bradshaw, H.
 submitted to the EMBL Data Library, February 1997
 A:Description: The sequence of C. elegans cosmid K11H12.
 A:Reference number: Z21526
 A:Accession: T34441
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA

A:Residues: 1-397 <BRA>
 A:Cross-references: EMBL:U88168; PIDN:AAC24400.1; GSPDB:GN00022; CESP:K11H12.3
 A:Experimental source: strain Bristol N2; clone K11H12
 C:Genetics:
 A:Gene: CESP:K11H12.3
 A:Map position: 4
 A:introns: 39/3; 68/2; 118/1; 206/2; 280/3

Query Match 81.6%; Score 31; DB 2; Length 397;
 Best Local Similarity 75.0%; Pred. No. 75;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 SIINFEXL 8
 |||||
 |||||
 Db 190 SIINFEXI 197

RESULT 15
 T42923
 infected cell protein - ateline herpesvirus 3 (strain 73)
 C:Species: ateline herpesvirus 3
 A:Variety: strain 73
 C:Date: 21-Jan-2000 #sequence revision 21-Jan-2000 #text_change 05-May-2000
 C:Accession: T42923
 R:Albrecht, J.C.; Fleckenstein, B.
 submitted to the EMBL Data Library, August 1998
 A:Description: Primary structure of the herpesvirus ateles genome.
 A:Reference number: Z22274
 A:Accession: T42923
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-680 <ALB>
 A:Cross-references: EMBL:AF083424; PIDN:AAC5539.1
 A:Experimental source: strain 73
 C:Genetics:
 A:Note: orf07
 C:Superfamily: herpesvirus infected cell protein ICP18.5

Query Match 81.6%; Score 31; DB 2; Length 680;
 Best Local Similarity 71.4%; Pred. No. 1.4e+02;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 SIINFEX 7
 |||||
 |||||
 Db 618 SVLNFEK 624

Search completed: January 30, 2004, 07:09:46
 Job time : 23 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 30, 2004, 07:06:27 ; Search time 12 Seconds
(without alignments)
31.351 Million cell updates/sec

Title: SEQ10
Perfect score: 38
Sequence: 1 sinife1 8

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues
Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_41.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	38	100.0	385	1 OVAL_CHICK	P01012 gallus galli
2	34	89.5	627	1 RNN3_YEAST	P36070 saccharomyc
3	33	86.8	382	1 OVAL_COTUJA	P19104 coturnix co
4	32	84.2	267	1 RMO9_HUMAN	O91472 homo sapien
5	31	81.6	242	1 TPIS_MYCEL	P48779 mycoplasma
6	31	81.6	244	1 YCAD_CYARA	P48334 cyamophila
7	31	81.6	304	1 MTH5_HAEIN	P45000 haemophilus
8	30	78.9	231	1 RLI_BUCAP	O8K667 buchnera ap
9	30	78.9	232	1 OVAX_CHICK	P01013 gallus galli
10	30	78.9	349	1 Y567_HELPJ	O92174 heliobacte
11	30	78.9	364	1 SERC_DROME	O92174 heliobacte
12	30	78.9	527	1 PRCK_FUSNN	O8E112 fusobacteri
13	30	78.9	558	1 MNT2_YEAST	P53059 saccharomyc
14	30	78.9	842	1 DRP3_THEMA	O92174 heliobacte
15	30	78.9	1163	1 SBCC_CLOAB	O971K1 clostridium
16	30	78.9	2469	1 TEGU_HSVSA	O01056 herpesvirus
17	30	78.9	4092	1 DVHC_YEAST	P34622 saccharomyc
18	29	76.3	143	1 PRIB_BACSU	P35137 bacillus su
19	29	76.3	237	1 YG56_METUA	O59050 methanococc
20	29	76.3	394	1 CC91_YEAST	P41313 saccharomyc
21	29	76.3	445	1 RPN5_YEAST	O12250 saccharomyc
22	29	76.3	453	1 EX7L_RICPR	O92174 heliobacte
23	29	76.3	482	1 V232_ROMPY	O92174 heliobacte
24	29	76.3	489	1 T283_STPAU	O92174 heliobacte
25	29	76.3	749	1 PCRA_LEUCI	O92174 heliobacte
26	29	76.3	1002	1 HP63_MOUSE	O92174 heliobacte
27	29	76.3	1233	1 SMCI_SCHPO	O92174 heliobacte
28	29	76.3	1234	1 YNKS_CAEEL	P34578 caenorhabdi
29	29	76.3	2054	1 YCR2_PINTA	P41653 pinus thunb
30	28	73.7	102	1 CTT1_ORISA	P09229 oxyza sativ
31	28	73.7	124	1 PA25_AGRKP	O42189 aglystodon
32	28	73.7	177	1 YASO_METUA	O58450 methanococc
33	28	73.7	214	1 PYRE_PASWU	O9CJW4 pasteurrella

34	28	73.7	222	1 BID2_YERPE	O9agd4 yersinia pe
35	28	73.7	240	1 YDED_SCHPO	O10446 schizosacch
36	28	73.7	242	1 TPIS_MYCH	P50920 mycoplasma
37	28	73.7	269	1 HIS9_LACLA	O02150 lactococcus
38	28	73.7	278	1 TNP6_RAT	P35940 rattus norv
39	28	73.7	279	1 TNP6_MOUSE	P41047 mus musculu
40	28	73.7	306	1 COA4_STRP3	O8B7C7 streptococc
41	28	73.7	306	1 COA4_STRP3	O8B0V9 streptococc
42	28	73.7	306	1 COA4_STRP3	O992h1 streptococc
43	28	73.7	309	1 LDH2_LACPL	P59390 lactobacill
44	28	73.7	315	1 ACC1_LYCES	P05116 lycopersico
45	28	73.7	316	1 ACC2_LYCES	P07920 lycopersico

ALIGNMENTS

RESULT 1
ID OVAL_CHICK STANDARD; PRT; 385 AA.
AC P01012;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Ovalbumin (Plakalbumin) (Allergen Gal d 2) (Gal d II).
OS Gallus gallus (Chicken).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;
OC Gallus.
OK NCBI_TaxID=9031;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=82069038; PubMed=6272839;
RA Brownlee G.G., O'Malley B.W., Catterall J.F., Dugaiczky A., Staden R.,
RA Woo S.L.C., Beattie W.G., Catterall J.F., Dugaiczky A., Staden R.,
RT "Complete nucleotide sequence of the chicken chromosomal ovalbumin
RT gene and its biological significance.",
RL Biochemistry 20:6437-6446(1981).
[2]
RP SEQUENCE FROM N.A.
RX MEDLINE=78199842; PubMed=661981;
RA McReynolds L., O'Malley B.W., Nisbet A.D., Fothergill J.E., Givol D.,
RA Fields S., Robertson M., Brownlee G.G.,
RT "Sequence of chicken ovalbumin mRNA.",
RL Nature 273:723-728(1978).
[3]
RP SEQUENCE FROM N.A.
RX MEDLINE=79010682; PubMed=692731;
RA Catterall J.F., O'Malley B.W., Robertson M.A., Staden R.,
RA Tanaka Y., Brownlee G.G.,
RT "Nucleotide sequence homology at 12 intron-exon junctions in the
RT chick ovalbumin gene.",
RL Nature 275:510-513(1978).
[4]
RP SEQUENCE OF 1-35.
RX MEDLINE=78116057; PubMed=272676;
RA Palmiter R.D., Gagnon J., Walsh K.A.,
RT "Ovalbumin: a secreted protein without a transient hydrophobic leader
RT sequence.",
RL Proc. Natl. Acad. Sci. U.S.A. 75:94-98(1978).
[5]
RP SEQUENCE OF 1-16.
RX MEDLINE=79186958; PubMed=751625;
RA Thompson E.O.P., Fisher W.K.,
RT "A correction and extension of the acetylated amino terminal sequence
RT of ovalbumin.",
RL Aust. J. Biol. Sci. 31:443-446(1978).
[6]
RP SEQUENCE OF 5-16; 29-35; 60-78; 115-123; 366-373 AND 379-385.
RX MEDLINE=79186957; PubMed=751624;
RA Thompson E.O.P., Fisher W.K.,
RT "Amino acid sequences containing half-cysteine residues in ovalbumin.",
RL Aust. J. Biol. Sci. 31:433-442(1978).

[7]
 RN N-TERMINUS ACETYLATION, AND PHOSPHORYLATION OF SER-68; SER-236 AND
 RP SER-240.
 RX MEDLINE=22056091; PubMed=12060738;
 RA MacCoss W.J., McDonald W.H., Saraf A., Sadygov R., Clark J.M.,
 RA Taabto J.J., Gould K.L., Wolters D., Washburn M., Weiss A., Clark J.I.,
 RA Yates J.R. III;
 RT "Shotgun identification of protein modifications from protein
 RT complexes and lens tissue.";
 RT Proc. Natl. Acad. Sci. U.S.A. 99:7900-7905(2002).
 RN [8]
 RP X-RAY CRYSTALLOGRAPHY (1.95 ANGSTROMS).
 RX MEDLINE=90370102; PubMed=2395463;
 RA Stein P.E., Leslie A.G.W., Finch J.T., Turnell W.G., McLaughlin P.J.,
 RA Carrell R.W.;
 RT "Crystal structure of ovalbumin as a model for the reactive centre of
 RT serpin.";
 RT Nature 347:99-102(1990).
 RN [9]
 RP X-RAY CRYSTALLOGRAPHY (1.95 ANGSTROMS).
 RX MEDLINE=92046044; PubMed=1942038;
 RA Stein P.E., Leslie A.G.W., Finch J.T., Carrell R.W.;
 RT "Crystal structure of uncleaved ovalbumin at 1.95-A resolution.";
 RT J. Mol. Biol. 221:941-959(1991).
 RN [10]
 RP X-RAY CRYSTALLOGRAPHY (2.8 ANGSTROMS).
 RX MEDLINE=90278960; PubMed=2352279;
 RA Wright H.T., Qian H.X., Huber R.;
 RT "Crystal structure of plakalbumin, a proteolytically nicked form of
 RT ovalbumin. Its relationship to the structure of cleaved alpha-1-
 RT proteinase inhibitor.";
 RT J. Mol. Biol. 213:513-528(1990).
 RN [11]
 RP REVIEW.
 RX MEDLINE=21312433; PubMed=11419711;
 RA Huntington J.A., Stein P.E.;
 RT "Structure and properties of ovalbumin.";
 RT J. Chromatogr. B 756:189-198(2001).
 CC -1- FUNCTION: Not known.
 CC -1- SUBCELLULAR LOCATION: Secreted.
 CC -1- TISSUE SPECIFICITY: Major protein of egg white.
 CC -1- SIMILARITY: BELONGS TO THE SERPIN FAMILY. OV-SERPIN SUBFAMILY.
 CC -1- DATABASE: NAME=Worthington enzyme manual;
 CC WWW="http://www.worthington-biochem.com/manual/O/OA.html".
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
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 CC modified and this statement is not removed. Usage by and for commercial
 CC entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 CC or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; J00895; AAB5956.1; -;
 DR EMBL; V00438; CAA23716.1; -;
 DR EMBL; V00383; CAA23682.1; -;
 DR EMBL; M34352; AAA48998.1; -;
 DR EMBL; M34346; AAA48998.1; JOINED.
 DR EMBL; M34347; AAA48998.1; JOINED.
 DR EMBL; M34348; AAA48998.1; JOINED.
 DR EMBL; M34349; AAA48998.1; JOINED.
 DR EMBL; M34350; AAA48998.1; JOINED.
 DR EMBL; M34351; AAA48998.1; JOINED.
 DR FIR; A90455; ORCH.
 DR PDB; 1OVA; 15-JUL-92.
 DR PDB; 1JTI; 18-DEC-02.
 DR PDB; 1VAC; 20-JUN-96.
 DR GlycoSuiteDB; P01012; -;
 DR InterPro; IPR000215; Serpin.
 DR Pfam; PF00079; Serpin.1.
 DR SMART; SM00093; SERPIN; 1.
 DR PROSITE; PS00284; SERPIN; 1.
 KW Serpin; Acetylation; Phosphorylation; Glycoprotein; 3D-structure;

KW	Allergen.	0	0	
FT	INIT MET	1	1	
FT	MOD RES	68	68	ACETYLATION.
FT	MOD RES	73	120	PHOSPHORYLATION.
FT	DISULFID	236	236	PHOSPHORYLATION.
FT	MOD RES	240	240	PHOSPHORYLATION.
FT	CARBOHYD	292	292	N-LINKED (GLCNAC. . .).
FT	MOD RES	344	344	PHOSPHORYLATION.
FT	ACT SITE	352	353	REACTIVE BOND HOMOLOG.
FT	VARIANT	311	311	N -> D (IN A MINOR COMPONENT).
FT	CONFLICT	187	187	A -> T (IN REF. 2).
FT	HELIIX	3	21	
FT	TURND	23	29	
FT	STRAND	27	24	
FT	HELIIX	31	43	
FT	TURND	44	44	
FT	HELIIX	47	57	
FT	TURND	58	58	
FT	TURND	60	61	
FT	HELIIX	63	64	
FT	HELIIX	67	70	
FT	TURND	71	75	
FT	TURND	79	80	
FT	HELIIX	81	91	
FT	STRAND	97	108	
FT	TURND	109	110	
FT	STRAND	113	113	
FT	HELIIX	115	124	
FT	STRAND	129	132	
FT	TURND	135	137	
FT	HELIIX	138	152	
FT	TURND	153	155	
FT	TURND	163	164	
FT	TURND	168	169	
FT	STRAND	172	182	
FT	STRAND	184	186	
FT	HELIIX	190	192	
FT	STRAND	194	199	
FT	STRAND	205	222	
FT	TURND	223	226	
FT	STRAND	227	234	
FT	TURND	235	236	
FT	STRAND	239	246	
FT	TURND	249	251	
FT	HELIIX	252	258	
FT	HELIIX	261	267	
FT	TURND	268	268	
FT	TURND	270	272	
FT	STRAND	274	283	
FT	STRAND	285	292	
FT	HELIIX	293	299	
FT	TURND	300	301	
FT	HELIIX	304	306	
FT	TURND	308	309	
FT	TURND	313	315	
FT	STRAND	319	316	
FT	TURND	319	320	
FT	STRAND	325	334	
FT	STRAND	338	340	
FT	STRAND	357	359	
FT	STRAND	364	370	
FT	TURND	371	373	
FT	STRAND	376	382	
SQ	SEQUENCE	385 AA;	42750 MW;	52B4339642388FE3 CRC64;

Query Match
 Best Local Similarity 100.0%; Score 38; DB 1; Length 385;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 STINFEKL 8
 |||||
 Db 257 STINFEKL 264

RESULT 2

```

RN3_YEAST      STANDARD;      PRT;      627 AA.
ID_RN3_YEAST
AC P36070.
DT 01-JUN-1994 (Rel. 29, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE RNA polymerase I specific transcription initiation factor RN3.
GN RN3 OR YKL12W.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomyces.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A., AND CHARACTERIZATION.
RX MEDLINE=96324404; PubMed=8670901;
RY Yamamoto R.T., Nogai Y., Dodd J.A., Nomura M.;
RT "RN3 gene of Saccharomyces cerevisiae encodes an essential RNA
RT polymerase I transcription factor which interacts with the polymerase
RT independently of DNA template.";
RL EMBL J. 15:364-3973 (1996).
RN [2]
RP SEQUENCE FROM N.A.
RA Rad M.R., Xu G., Kirchrath L., Fritz C., Keuchel H., Hollenberg C.P.;
RL Submitted (MAR-1994) to the EMBL/Genbank/DBJ databases.
CC -1- FUNCTION: REQUIRED FOR EFFICIENT TRANSCRIPTION INITIATION BY RNA
CC POLYMERASE I. INTERACTS WITH POL I IN THE ABSENCE OF TEMPLATE DNA
CC AND STIMULATES RECRUITMENT OF POL I, BUT DOES NOT REMAIN AS PART
CC OF STABLE PRE-INITIATION COMPLEX.
CC -1- SUBUNIT: Monomer (Probable).
CC -1- SUBCELLULAR LOCATION: Nuclear (Potential).
CC -1- SIMILARITY: TO S.POMBE SPAC1866.11C AND C.ELEGANS C36B8.1.
CC
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CC -----
DR EMBL; Z71927; CAA96470.1; -
DR EMBL; Z28125; CAA81966.1; -
DR PIR; S37954; S37954.
DR TRANSPAC; T03565; -
DR SGD; S0001608; RN3.
DR Pfam; PF05327; RN3; 1.
KW Transcription regulation; Nuclear protein.
FT DOMAIN 252 259 POLY-ASP.
FT DOMAIN 267 274 POLY-ASP.
FT DOMAIN 277 280 POLY-ASP.
FT DOMAIN 546 549 POLY-ASN.
SQ SEQUENCE 627 AA; 72387 MW; A31E7386A987FDB CRC64;

Query Match      89.5%; Score 34; DB 1; Length 627;
Best Local Similarity 100.0%; Pred. No. 11;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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OS Coturnix coturnix japonica (Japanese quail).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;
OC Coturnix.
OX NCBI_TaxID=93934;
RN [1]
RP SEQUENCE FROM N.A.
RC TISUE=Oviduct;
RX MEDLINE=91016850; PubMed=2216734;
RA Mucha J., Klaidinyova V., Hanes J., Simuth J.;
RT "The sequence of Japanese quail ovalbumin cDNA.";
RL Nucleic Acids Res. 18:5553-5553 (1990).
CC -1- SIMILARITY: BELONGS TO THE SERPIN FAMILY. OV-SERPIN SUBFAMILY.
CC
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CC -----
DR EMBL; X53964; CAA37916.1; -
DR PIR; S11433; S11433.
DR HSSP; P01012; IOVA.
DR InterPro; IPR000215; Serpin.
DR Pfam; PF00079; serpin; 1.
DR SMART; SM00093; SERPIN; 1.
DR PROSITE; PS00284; SERPIN; 1.
KW Serpin; Glycoprotein.
FT INIT MET 0
FT DISULFID 73 120 BY SIMILARITY.
FT CARBOHYD 292 292 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT ACT SITE 311 311 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT ACT SITE 352 353 REACTIVE BOND HOMOLOG.
SQ SEQUENCE 362 AA; 42108 MW; A4BB59A1BAE8F316 CRC64;

Query Match      86.8%; Score 33; DB 1; Length 382;
Best Local Similarity 87.5%; Pred. No. 11;
Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

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OY 1 SIINFEX 7
DB 118 SIINFEX 124

RESULT 3
OVAL_COTUA      STANDARD;      PRT;      382 AA.
ID_OVAL_COTUA
AC P19104.
DT 01-NOV-1990 (Rel. 16, Created)
DT 01-NOV-1990 (Rel. 16, Last sequence update)
DT 01-APR-1993 (Rel. 25, Last annotation update)
DE Ovalbumin.

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OY 1 SIINFEX 8
DB 257 SIINFEX 264

RESULT 4
RM09_HUMAN      STANDARD;      PRT;      267 AA.
ID_RM09_HUMAN
AC O98YD2; O9BSW8;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE 60S ribosomal protein l9, mitochondrial precursor (L9mt).
GN MRP19.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21293042; PubMed=11279069;
RA Suzuki T., Terasaki M., Takemoto-Hori C., Hanada T., Ueda T., Wada A.,
RT "Structural compensation for the deficit of rRNA with proteins in the
RT mammalian mitochondrial ribosome. Systematic analysis of protein
RT components of the large ribosomal subunit from mammalian
RT mitochondria.";
RL J. Biol. Chem. 276:21724-21736 (2001).
RN [2]
RP SEQUENCE FROM N.A.
RC TISUE=Lung;
RX MEDLINE=22388257; PubMed=12477932;

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RA Strausberg R.L., Feigold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shemmer C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Bueger K.H., Scheffer C.F., Bat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diachenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Schetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Locquellano N.A., Peters G.J., Abramson R.D., Mullaly S.J.,
RA Bosnak S.A., McEwen P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Morley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalón D.K., Mizny D.M., Sodergren E.U., Lu X., Gibbs R.A.,
RA Fahy J., Helton E., Keltman M., Madan A., Rodrigues S., Sanchez A.,
RA Blakesley R.W., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Rodriguez A.C., Touchman J.W., Green E.D., Dickson M.C.,
RA Butterfield V.S.N., Krzywinski M.I., Skalska U., Smallos D.E.,
RA Schermer A., Schein J.E., Jones S.J.M., Maira M.A.,
RT "Generation and initial analysis of more than 15,000 full-length
RT human and mouse cDNA sequences."
RT Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
CC -1- SUBCELLULAR LOCATION: Mitochondrial.
CC -1- SIMILARITY: BELONGS TO THE L9P FAMILY OF RIBOSOMAL PROTEINS.
CC -----
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CC -----
DR EMBL; AB049636; BAB40841.1; -
DR EMBL; BC004517; AAH04517.1; -
DR Genew; HGNC:14277; MRPL9.
DR InterPro; IPR000244; Ribosomal_L9.
DR Pfam; PF01281; Ribosomal_L9_N; 1.
DR KMW Ribosomal protein; Mitochondrion; Transic peptide.
FT TRANSLIT 1 267 MITOCHONDRION (POTENTIAL).
FT CHAIN ? 267 60S RIBOSOMAL PROTEIN L9.
FT CONFLICT 210 210 A -> E (TN REF. 2).
SQ SEQUENCE 267 AA; 30185 MW; 346C254220FDB4 CRC64;

Query Match
Best Local Similarity 84.2%; Score 32; DB 1; Length 267;
Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEK 7
ID 1:|||||
AC P48779; STANDARD; PRT; 242 AA.
DB 236 SVINFEK 242

RESULT 5
TPIS MYCFL STANDARD; PRT; 242 AA.
AC P48779;
DT 01-FEB-1996 (Rel. 33, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Triosephosphate isomerase (EC 5.3.1.1) (TIM).
GN TPIA OR TPI.
OS Mycoplasma flocculare.
OC Bacteria; Firmicutes; Mollicutes; Mycoplasmataceae; Mycoplasma.
OX NCBI_TaxID=2128;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 27399;
RA Xiang H., McIntosh M.A.;
RL Submitted (JAN-1995) to the EMBL/GenBank/DDA databases.
CC -1- CATALYTIC ACTIVITY: D-glyceraldehyde 3-phosphate = glyceralone
CC phosphate.
CC -1- PATHWAY: Plays an important role in several metabolic pathways.
CC -1- SUBUNIT: Homodimer (By similarity).
CC -1- SIMILARITY: BELONGS TO THE TRIOSEPHOSPHATE ISOMERASE FAMILY.

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CC -----
DR EMBL; U20509; AAA62167.1; -
DR HSSP; P00943; 2BTM.
DR HAMAP; MF_00147; 1.
DR InterPro; IPR000652; Triophos_ismrse.
DR Pfam; PF00121; TIM; 1.
DR ProDom; PD001005; Triophos_ismrse; 1.
DR TIGRPFAMs; TIGR00419; tim; 1.
DR PROSITE; PS00171; TIM; 1.
DR Isomerase; Glycolysis; Gluconeogenesis; Fatty acid biosynthesis;
KM Pentose shunt.
FT ACT SITE 98 98 BY SIMILARITY.
FT ACT SITE 167 167 BY SIMILARITY.
SQ SEQUENCE 242 AA; 26969 MW; B1E560E2DA18F41 CRC64;

Query Match
Best Local Similarity 81.6%; Score 31; DB 1; Length 242;
Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEK 7
ID 1:|||||
DB 155 SVINFEK 161

RESULT 6
YXCD CYAPA
ID YXCD CYAPA STANDARD; PRT; 244 AA.
AC P48334;
DT 01-FEB-1996 (Rel. 33, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Probable ABC transporter ATP-binding protein in yef23-apef intergenic
DE region (ORF244).
OS Cyanophora paradoxa.
OG Cyanella.
OC Eukaryota; Glaucocystophyceae; Cyanophoraceae; Cyanophora.
OX NCBI_TaxID=2762;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=UTEX LB 555 / Pringsheim;
RA Stirewalt V.L., Michalowski C.B., Loeffelhardt W., Bohnert H.J.,
RA Bryant D.A.;
RT "Nucleotide sequence of the cyanella DNA from Cyanophora paradoxa.";
RL Plant Mol. Biol. Rep. 13:327-332(1995).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=UTEX LB 555 / Pringsheim;
RA Loeffelhardt W., Stirewalt V.L., Michalowski C.B., Annarella M.,
RA Farley J.V., Schlueter W.M., Chung S., Neumann-Spallart C.,
RA Steiner J.M., Jakowitsch J., Bohnert H.J., Bryant D.A.;
RT "The complete sequence of the cyanella genome of Cyanophora paradoxa:
RT the genetic complexity of a primitive plastid.";
RL (in) Schenk H.E.A., Herrmann R., Jeon K.W., Mueller N.E.,
RL Schwemmler W. (eds.);
RL Eukaryotism and Symbiosis, pp.40-48, Springer-Verlag, Heidelberg
RL (1997).
CC -1- SIMILARITY: Belongs to the ABC transporter family.
CC -----
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CC -----
CC EMBL; U30821; AAA81304.1; -
CC PIR; T06961; T06961.
CC InterPro; IPR003593; AAA_Arase.
CC InterPro; IPR003439; ABC_transporter.
CC Pfam; PF00005; ABC_tran; 1.
CC ProDom; PD000006; ABC_transporter; 1.
CC SMART; SM00382; AAA; 1.
CC PROSITE; PS00211; ABC_TRANSPORTER_1; 1.
CC PROSITE; PS00893; ABC_TRANSPORTER_2; 1.
CC Hypothetical protein; ATP-binding; Transport; Cyanelle.
CC NP_BIND 41 48 ATP (POTENTIAL).
CC SEQUENCE 244 AA; 27747 MW; 4CSB357F9C55D3B CRC64;

Query Match
Best Local Similarity 81.6%; Score 31; DB 1; Length 244;
Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

OY 1 STINFEKT 8
DB 204 STINFEKT 211

RESULT 7
MTH5_HAEIN STANDARD; PRT; 304 AA.
AC P45000;
DT 01-NOV-1995 (Rel. 32, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DE Modification methylase Hindv (EC 2.1.1.73) (Cytosine-specific
DE methyltransferase Hindv) (M.Hindv).
GN HINDVM OR H11041.
OS Haemophilus influenzae.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Pasteurellales;
OC Pasteurellaceae; Haemophilus.
OX NCBI_TaxID=727;
[1]
RP SEQUENCE FROM N.A.
RC STRAIN=RD / KW20 / ATCC 51907;
RX MEDLINE=95350630; PubMed=7542800;
RA Fleischmann R.D., Adams M.D., White O., Clayton R.A., Kirkness E.F.,
RA Kerlavage A.R., Bult C.J., Tomb J.-F., Dougherty B.A., Merrick J.M.,
RA McKenney K., Sutton G., Fitzhugh W., Fields C.A., Gocayne J.D.,
RA Scott J.D., Shiley R., Liu L.-I., Glodek A., Kelley J.M.,
RA Weisman J.F., Phillips C.A., Spriggs T., Hedblom E., Cotton M.D.,
RA Usterback T.R., Hanna M.C., Nguyen D.T., Saudek D.M., Brandon R.C.,
RA Fine L.D., Fritchman J.L., Fuhmann J.L., Geoghegan N.S.M.,
RA Gnehm C.L., McDonald L.A., Small K.V., Fraser C.M., Smith H.O.,
RA Venter J.C.;
RA "Whole-genome random sequencing and assembly of Haemophilus influenzae
RT Rd.";
RL Science 269:496-512 (1995).
CC -1- FUNCTION: THIS METHYLASE RECOGNIZES THE DOUBLE-STRANDED SEQUENCE
CC GRCGVC, CAUSES SPECIFIC METHYLATION ON C-? ON BOTH STRANDS, AND
CC PROTECTS THE DNA FROM CLEAVAGE BY THE HINDV ENDONULEASE.
CC -1- CATALYTIC ACTIVITY: S-adenosyl-L-methionine + DNA cytosine = S-
CC adenosyl-L-homocysteine + DNA 5-methylcytosine.
CC -1- SIMILARITY: BELONGS TO THE C5-METHYLTRANSFERASE FAMILY.
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CC -----
CC EMBL; U32784; AAC22700.1; -
CC PIR; C64109; C64109.
CC HSSP; O14717; 1G55.
CC REBASE; 3574; M.Hindv.
CC TIGR; H11041; -

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DR InterPro; IPR001525; C5_DNA_meth.
DR Pfam; PF00145; DNA_methylase; 1.
DR PRINTS; PRO0105; CSMETTRFRASE.
DR TIGRPFAM; TIGR00675; dcm; 1.
DR PROSITE; PS00094; C5_MTASE_1; 1.
DR PROSITE; PS00095; C5_MTASE_2; 1.
DR Hypothetical protein; Transferase; Methyltransferase;
DR Restriction system; Complete proteome.
FT ACT SITE 75 75
BY SIMILARITY.
SQ SEQUENCE 304 AA; 34365 MW; 03DA1EAB27C84BBD CRC64;

Query Match
Best Local Similarity 81.6%; Score 31; DB 1; Length 304;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 1 STINFEKT 7
DB 291 STINFEKT 297

RESULT 8
RL1_BUCAP STANDARD; PRT; 231 AA.
ID RL1_BUCAP
AC Q8KA67;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE 50S ribosomal protein L1.
GN RPLA OR BUSG038.
OS Buchnera aphidicola (subsp. Schizaphis graminum).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Buchnera.
OX NCBI_TaxID=98794;
[1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22084549; PubMed=12089438;
RA Tamas I., Klaesson L., Canbaeck B., Naeslund A.K., Eriksson A.-S.,
RA Wernegreen J.J., Sandstroem J.P., Moran N.A., Andersson S.G.E.;
RT "50 million years of genomic stasis in endosymbiotic bacteria.";
RL Science 296:2376-2379 (2002).
CC -1- FUNCTION: THIS PROTEIN BINDS DIRECTLY TO 23S RIBOSOMAL RNA AND IS
CC LOCATED IN THE NEIGHBORHOOD OF THE SITE WHERE ELONGATION FACTOR TU
CC IS BOUND TO THE RIBOSOME (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE L1P FAMILY OF RIBOSOMAL PROTEINS.
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CC -----
CC EMBL; AE014080; AAM67609.1; -
CC InterPro; IPR005878; L1_bact chl.
CC InterPro; IPR002143; Ribosomal_L1.
CC Pfam; PF00687; Ribosomal_L1; 1.
CC ProDom; PD001314; Ribosomal_L1; 1.
CC TIGRPFAM; TIGR01169; rplA_bact; 1.
CC PROSITE; PS01199; RIBOSOMAL_L1; 1.
CC Ribosomal protein; rRNA-binding; Complete proteome.
SQ SEQUENCE 231 AA; 25609 MW; 4D334DACF932A3C2 CRC64;

Query Match
Best Local Similarity 78.9%; Score 30; DB 1; Length 231;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 INFEKT 8
DB 15 INFEKT 20

RESULT 9

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OVAX CHICK STANDARD; PRT; 232 AA.
 ID OVAX CHICK
 AC Q92LR4;
 DT 21-JUL-1986 (Rel. 01, Created)
 DT 21-JUL-1986 (Rel. 01, Last sequence update)
 DT 01-APR-1993 (Rel. 25, Last annotation update)
 DE Gene X protein (Ovalbumin-related) (Fragment).
 GN X.
 OS Gallus gallus (Chicken).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Archaeosuria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae;
 OC Gallus.
 OX NCBI_TaxID=9031;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=81022623; PubMed=7418002;
 RA Helling R., Petrin F., Gannon F., Mandel J.L., Chambon P.;
 RT "The ovalbumin gene family: structure of the X gene and evolution of
 duplicated split genes."
 RL Cell 20:625-637(1980).
 CC -1- SIMILARITY: BELONGS TO THE SERPIN FAMILY. OY-SERPIN SUBFAMILY.
 CC -----
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 CC -----
 CC DR EMBL; J00920; AAA68881.1; -
 DR EMBL; J00918; AAA68881.1; JOINED.
 DR EMBL; J00919; AAA68881.1; JOINED.
 DR EMBL; V00385; CAA23683.1; -
 DR EMBL; V00386; CAA23684.1; -
 DR EMBL; V00387; CAA23685.1; -
 DR PIR; A01243; DXCH.
 DR HSSP; P01012; IOVA.
 DR InterPro: IPR000215; Serpin.
 DR Pfam; PF00079; serpin.1.
 DR SMART; SM00093; SERPIN; 1.
 DR PROSITE; PS00284; SERPIN; 1.
 KW Serpin.
 FT NON TER
 SQ SEQUENCE 232 AA; 26291 MW; 6B5B86EC4D3B195 CRC64;
 Query Match 78.9%; Score 30; DB 1; Length 232;
 Best Local Similarity 100.0%; Pred. NO. 29;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 3 INFECL 8
 DB 104 INFECL 109
 RESULT 10
 Y567_HELPJ STANDARD; PRT; 349 AA.
 AC Q92LR4;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Hypothetical protein JHP0514.
 GN JHP0514.
 OS Helicobacter pylori J99 (Campylobacter pylori J99).
 OC Bacteria; Proteobacteria; Epsilonproteobacteria; Campylobacterales;
 OC Helicobacteriaceae; Helicobacter.
 OX NCBI_TaxID=85963;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=99120557; PubMed=9923662;
 RA Alm R.A., Ling L.-S.L., Moir D.T., King B.L., Brown E.D., Doig P.C.,
 RA Smith D.R., Noonan B., Guild B.C., deJonge B.L., Carmel G.,

RA Tummino P.J., Caruso A., Uria-Nickelsen M., Mills D.M., Ives C.,
 RA Gibson R., Merberg D., Mills S.D., Jiang Q., Taylor D.E., Vovis G.F.,
 RA Trust T.J.;
 RT "Genomic sequence comparison of two unrelated isolates of the human
 RT gastric pathogen Helicobacter pylori."
 RL Nature 397:176-180(1999).
 CC -1- SUBCELLULAR LOCATION: Integral membrane protein (potential).
 CC -1- SIMILARITY: BELONGS TO THE UPF0118 (PERM) FAMILY.
 CC -----
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 CC -----
 CC DR EMBL; AE001484; AAD06090.1; -
 DR PIR; H71923; H71923.
 DR InterPro: IPR002549; UPF0118.
 DR Pfam; PF01594; UPF0118; 1.
 KW Hypothetical protein; Transmembrane; Complete proteome.
 FT TRANSMEM 6 26 POTENTIAL.
 FT TRANSMEM 27 47 POTENTIAL.
 FT TRANSMEM 56 76 POTENTIAL.
 FT TRANSMEM 143 163 POTENTIAL.
 FT TRANSMEM 195 215 POTENTIAL.
 FT TRANSMEM 224 244 POTENTIAL.
 FT TRANSMEM 258 278 POTENTIAL.
 FT TRANSMEM 300 320 POTENTIAL.
 SQ SEQUENCE 349 AA; 39804 MW; A1846D48CB3A86F CRC64;
 Query Match 78.9%; Score 30; DB 1; Length 349;
 Best Local Similarity 100.0%; Pred. NO. 44;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 3 INFECL 8
 DB 85 INFECL 90
 RESULT 11
 SERC DROME STANDARD; PRT; 364 AA.
 AC Q9VAN0;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Probable phosphoserine aminotransferase (EC 2.6.1.52) (PSAT).
 GN EST3:39C108 OR CG11899.
 GN Drosophila melanogaster (Fruit fly).
 OS Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
 OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 OX NCBI_TaxID=7227;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Berkeley;
 RX MEDLINE=20196006; PubMed=10731132;
 RA Adams M.D., Celinker S.E., Holt R.A., Evans C.A., Gocayne J.D.,
 RA Amanatides P.G., Scherer S.E., Li P.W., Hoskins R.A., Galie R.F.,
 RA George R.A., Lewis S.E., Richards S., Ashburner M., Henderson S.N.,
 RA Sutton R.G., Wortman J.R., Yandell M.D., Zhang Q., Chen L.X.,
 RA Brandon R.C., Rogers Y.-H.C., Blake J.R.G., Chapple M., Pfeiffer B.D.,
 RA Wan K.H., Doyle C., Baxter E.G., Helt G., Nelson C.R., Miklos G.L.G.,
 RA Abril J.F., Agbayani A., An H.-J., Andrews-Pfannkoch C., Baldwin D.,
 RA Ballew R.M., Basu A., Baxendale J., Bayraktoglu L., Beasley E.M.,
 RA Beeson K.Y., Benos P.V., Bertan B.P., Bhandari D., Bolshakov S.,
 RA Borokova D., Botchan M.R., Bouck J., Brockstein P., Brotlier P.,
 RA Burtis K.C., Busam D.A., Butler H., Cadieu L.B., Davies P.,
 RA Cherry J.M., Cawley S., Dahlke C., Davenport L.B., Davies P.,
 RA de Pablos B., Delcher A., Deng Z., Mays A.D., Dew I., Dietz S.M.,
 RA Dodson K., Doup L.E., Downes M., Dugan-Rocha S., Dunkov B.C., Dunn P.,

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RA Durbin K.J., Evangelista C.C., Ferraz C., Ferriera S., Fleischmann W.,
RA Foster C., Garbalian A.E., Garg N.S., Gelbart W.M., Glaeser K.,
RA Glodex A., Gong F., Gorrell J.H., Gu Z., Guan P., Harris M.,
RA Harris N.L., Harvey D., Heiman T.J., Hernandez J.R., Houck J.,
RA Hostin D., Houston K.A., Howland T.J., Wei M.-H., Ibegwam C.,
RA Jatali M., Katush F., Karpen G.H., Ke Z., Kennison J.A., Ketchum K.A.,
RA Kimmel B.E., Kodira C.D., Kraft C., Kravitz S., Kulp D., Lai Z.,
RA Laeko P., Lei Y., Levitsky A.A., Li J., Li Z., Liang Y., Lin X.,
RA Liu X., Matzel B., McIntosh T.C., McLeod M.P., McPherson D.,
RA Melnikov G., Milphina N.V., Mobarry C., Morris J., Moshrefi A.,
RA Mount S.M., Moy M., Murphy B., Murphy L., Muzny D.M., Nelson D.L.,
RA Nelson D.R., Nelson K.A., Nixon K., Nusskern D.R., Paclet J.M.,
RA Palazzolo M., Pittman G.S., Pan S., Pollard J., Puri V., Reese M.G.,
RA Reibert K., Remington K., Saunders R.D.C., Scheeler F., Shen H.,
RA Shue B.C., Siden-Kiamos I., Simpson M., Skupski M.P., Smith T.,
RA Spletter E., Spradling A.C., Stapleton M., Strong R., Sun E.,
RA Svirskas R., Tector C., Turner R., Venter E., Wang A.H., Wang X.,
RA Wang Z.-Y., Wasserman D.A., Weinstein G.M., Weissbach J.,
RA Williams S.M., Woodage T., Worley K.C., Wu D., Yang S., Yao Q.A.,
RA Ye J., Yeh R.-F., Zaveri J.S., Zhan M., Zhang G., Zhao Q., Zheng L.,
RA Zheng X.H., Zhong F.N., Zhong W., Zhou X., Zhu S., Zhu X., Smith H.O.,
RA Gibbs R.A., Myers E.W., Rubin G.M., Venter J.C.;
RT "The genome sequence of Drosophila melanogaster.";
RL Science 287:2185-2195(2000).
CC -1- CATALYTIC ACTIVITY: O-phospho-L-serine + 2-oxoglutarate = 3-
CC phosphonoxypruvate + L-glutamate.
CC -1- COFACTOR: Pyridoxal phosphate.
CC -1- PATHWAY: REQUIRED BOTH IN MAJOR PHOSPHORYLATED PATHWAY OF SERINE
CC BIOSYNTHESIS AND IN THE BIOSYNTHESIS OF PYRIDOXINE.
CC -1- SIMILARITY: Belongs to class-V of pyridoxal-phosphate-dependent
CC aminotransferases.
-----
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-----
CC EMBL; AE003768; AAF56874.1; -.
CC DR HSPSP; P23721; IBDN.
CC DR FLYBASE; FBgn0014427; ESTS:39C10S.
CC DR InterPro; IPR000192; Aminotransf.
CC DR InterPro; IPR003248; Pser aminotransf.
CC DR Pfam; PF00266; aminotran_5; 1.
CC DR ProDom; PD001544; Pser_aminotransf; 1.
CC DR TIGRFAMs; TIGR01364; serC_1; 1.
CC DR PROSITE; PS00595; AA_TRANSFERS CLASS 5; 1.
CC KW Serine biosynthesis; Transferase; Aminotransferase;
CC PYRIDOXAL PHOSPHATE (BY SIMILARITY).
CC FT BINDING 194
CC SEQUENCE 364 AA; 39540 MW; DA6A4E2F5BD4DB74 CRC64;
SQ
Query Match 78.9%; Score 30; DB 1; Length 364;
Best Local Similarity 62.5%; Pred. No. 46;
Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
QY 1 SIINFEXL 8
DB 220 SIINFEXOM 227

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OS Fusobacterium nucleatum (subsp. nucleatum).
OC Bacteria; Fusobacteria; Fusobacteriales; Fusobacteriaceae;
OC Fusobacterium.
OX NCBI_TaxID=76856;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 25586;
RX MEDLINE=21886394; PubMed=11889109;
RA Kapralov V., Anderson I., Ivanova N., Reznik G., Los T., Lykidis A.,
RA Bhattacharya A., Bartman A., Gardner W., Greckin G., Zhu L.,
RA Larsen N., D'Souza M., Walunas T., Pusch G., Haselkorn R.,
RA Vasileva O., Chu L., Kogan Y., Chaga O., Goldsman E., Bernal A.,
RA Fomstein M., Kyridis N., Overbeek R.;
RT "Genome sequence and analysis of the oral bacterium Fusobacterium
RT nucleatum strain ATCC 25586."
RL J. Bacteriol. 184:2005-2018(2002).
CC -1- CATALYTIC ACTIVITY: ATP + oxaloacetate = ADP + phosphoenolpyruvate
CC + CO(2).
CC -1- PATHWAY: Rate-limiting gluconeogenic enzyme.
CC -1- SUBCELLULAR LOCATION: Cytoplasmic (by similarity).
CC -1- SIMILARITY: Belongs to the phosphoenolpyruvate carboxykinase (ATP)
CC family.
-----
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-----
CC EMBL; AE010616; AAL95316.1; -.
CC DR HAMAP; MF_00453; -.
CC DR InterPro; IPR001272; PEPCK_ATP.
CC DR Pfam; PF01293; PEPCK_ATP; 1.
CC DR ProDom; PD004723; PEPCK_ATP; 1.
CC DR TIGRFAMs; TIGR00224; pckA; 1.
CC DR PROSITE; PS00532; PEPCK_ATP; 1.
CC KW Gluconeogenesis; Lyase; Decarboxylase; ATP-binding; Complete proteome.
CC FT NP_BIND 230
CC SEQUENCE 527 AA; 59055 MW; 275849FDF254AC01 CRC64;
SQ
Query Match 78.9%; Score 30; DB 1; Length 527;
Best Local Similarity 100.0%; Pred. No. 66;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 2 IINFEX 7
DB 173 IINFEX 178

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RESULT 12
PCK_FUSNN STANDARD; PRT; 527 AA.
AC Q8RE12;
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Phosphoenolpyruvate carboxykinase [ATP] (EC 4.1.1.49) (PEP
DE carboxykinase) (Phosphoenolpyruvate carboxylase) (PEPCK).
GN PCKA OR FN1120.
RN [2]

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RESULT 13
MNT2 YEAST STANDARD; PRT; 558 AA.
AC P53059;
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Alpha-1,3-mannosyltransferase MNT2 (EC 2.4.1.-).
GN MNT2 OR YGL257C OR NR0558.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomyces.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=S288C / FY1679;
RX MEDLINE=97127827; PubMed=8972578;
RA Colasac E., Maillier E., Robineau S., Netter P.;
RT "Sequence of a 39/411 bp DNA fragment covering the left end of
RT chromosome VII of Saccharomyces cerevisiae."
RL Yeast 12:1555-1562(1996).
RN [2]

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RP CHARACTERIZATION.
RX MEDLINE=99453866; PubMed=10521541;
RA Romero P.A., Luseter M., Veronneau S., Sdicu A.M., Herscovics A.,
RA Bussey H.,
RT "Mnzp and Mutp of Saccharomyces cerevisiae are members of the Mnzp
RT family of alpha-1,3-mannosyltransferases responsible for adding the
RT terminal mannose residues of O-linked oligosaccharides."
RL Glycobiology 9:1045-1051(1999).
CC
CC -1- FUNCTION: Mannosyltransferase involved in adding the 4th and 5th
CC mannose residues of O-linked glycans.
CC
CC -1- PATHWAY: Glycosylation.
CC
CC -1- SUBCELLULAR LOCATION: Type II membrane protein. Golgi (Potential).
CC
CC -1- SIMILARITY: BELONGS TO THE MN1/MNT FAMILY.
CC
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CC -----
CC EMBL: X94357; CAA64130.1; -
CC EMBL: 272779; CAA96977.1; -
CC PIR: S61604; S61604.
CC SCD: S0003226; MNT2.
CC GO: GO:0000033; F:alpha-1,3-mannosyltransferase activity; IDA.
CC GO: GO:0006493; P:O-linked glycosylation; IDA.
CC Transferrase; Glycosyltransferase; Glycoprotein; Transmembrane;
CC Signal-anchor; Golgi stack.
CC TRANSMEM 1 27 CYTOPLASMIC (POTENTIAL).
CC FT DOMAIN 6 SIGNAL-ANCHOR (TYPE-II MEMBRANE PROTEIN)
CC FT TRANSMEM 1 27 (POTENTIAL).
CC FT DOMAIN 6 LUMENAL (POTENTIAL).
CC FT CARBOHYD 187 187 N-LINKED (GLYCNAC. . .) (POTENTIAL).
CC FT SEQUENCE 558 AA; 64852 MW; 3E58ED2B4E291B6 CRC64;
SQ
Query Match 78.9%; Score 30; DB 1; Length 558;
Best Local Similarity 100.0%; Pred. No. 70;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 INFEKL 8
DB 109 INFEKL 114

RESULT 14
DP3A_THEME STANDARD; PRT; 842 AA.
AC Q9ZHG4;
DT 30-MAY-2000 (Rel. 39, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE DNA polymerase III alpha subunit (EC 2.7.7.7).
GN DNAP OR TM0461.
OS Thermotoga maritima.
OC Bacteria; Thermotogae; Thermotogales; Thermotogaceae; Thermotoga.
OX NCBI_Taxid=2336;
RN (1)
RP SEQUENCE FROM N.A.
RX MEDLINE=99045593; PubMed=9826752;
RA Huang Y.P., Ito Y.,
RT "The hyperthermophilic bacterium Thermotoga maritima has two different
RT classes of family C DNA polymerases: evolutionary implications."
RL Nucleic Acids Res. 26:5300-5309(1998).
RN (2)
RP SEQUENCE FROM N.A.
RX STRAIN=MSB / DSM 3109;
RX MEDLINE=99287316; PubMed=10360571;
RA Nelson K.E., Clayton R.A., Gill S.R., Gwinn M.L., Dodson R.J.,
RA Haft D.H., Hickey E.K., Peterson J.D., Nelson W.C., Ketchum K.A.,
RA McDonald L., Uterback T.R., Malek J.A., Linher K.D., Garrett M.M.,
RA Stewart A.M., Cotton M.D., Pratt M.S., Phillips C.A., Richardson D.,

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RA Heidelberg J., Sutton G.G., Fleischmann R.D., Eisen J.A., White O.,
RA Salzberg S.L., Smith H.O., Venter J.C., Fraser C.M.;
RT "Evidence for lateral gene transfer between Archaea and Bacteria from
RT genome sequence of Thermotoga maritima."
RL Nature 399:323-329(1999).
CC
CC -1- FUNCTION: DNA POLYMERASE III IS A COMPLEX, MULTICHAIN ENZYME
CC RESPONSIBLE FOR MOST OF THE REPLICATIVE SYNTHESIS IN BACTERIA.
CC THIS DNA POLYMERASE ALSO EXHIBITS 3' TO 5' EXONUCLEASE ACTIVITY.
CC THE ALPHA CHAIN IS THE DNA POLYMERASE (BY SIMILARITY).
CC -1- CATALYTIC ACTIVITY: N deoxynucleoside triphosphate = N diphosphate
CC + [DNA] (N).
CC
CC -1- SUBUNIT: DNA polymerase III contains a core (composed of alpha,
CC epsilon and theta chains) that associates with a tau subunit. This
CC core dimerizes to form the POLIIT' complex. Politi' associates
CC with the gamma complex (composed of gamma, delta, delta', psi and
CC chi chains) and with the beta chain to form the complete DNA
CC polymerase III complex (By similarity).
CC
CC -1- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
CC
CC -1- SIMILARITY: BELONGS TO THE DNA POLYMERASE TYPE-C FAMILY. DNASE
CC SUBFAMILY.
CC
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CC -----
CC EMBL: AF063188; AAC80434.1; -
CC EMBL: AB001724; AAD35546.1; -
CC PIR: E72373; E72373.
CC TIGR: TM0461; -
CC DR Interpro: IPR003141; PNP_N.
CC DR Interpro: IPR004805; POLC_alpha.
CC DR SMART: SM00481; POLIITAC; 1.
CC DR TIGRFAMs: TIGR00594; polc; 1.
CC Transferrase; DNA-directed DNA polymerase; DNA replication;
CC Complete proteome.
CC KW Complete proteome. 377 I -> M (IN REF. 1).
CC FT CONFLICT 377
CC FT SEQUENCE 842 AA; 96499 MW; 19BB813C8085C277 CRC64;
SQ
Query Match 78.9%; Score 30; DB 1; Length 842;
Best Local Similarity 62.5%; Pred. No. 11e+02;
Matches 5; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 STINFEKL 8
DB 14 SVRFEKL 21

RESULT 15
SBCC_CLOAB STANDARD; PRT; 1163 AA.
ID SBCC_CLOAB
AC Q97FK1;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Nuclease bDCC subunit C.
GN SBCC OR CAC2736.
OS Clostridium acetobutylicum.
OC Bacteria; Firmicutes; Clostridia; Clostridiales; Clostridiaceae;
OC Clostridium.
OX NCBI_Taxid=1488;
RN (1)
RP SEQUENCE FROM N.A.
RX STRAIN=ATCC 824 / DSM 792 / VKM B-1787;
RX MEDLINE=2139325; PubMed=11466286;
RA Niessing U., Breton G., Omelchenko M.V., Makarova K.S., Zeng Q.,
RA Gibson R., Lee H.M., Dubois J., Qiu D., Hitti J., Wolf Y.I.,
RA Tatusov R.L., Sabathe F., Doucette-Stamm L., Soucaille P., Daly M.J.,
RA Bennett G.N., Koonin E.V., Smith D.R.;
RT "Genome sequence and comparative analysis of the solvent-producing

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RT bacterium Clostridium acetobutylicum.;
RL J. Bacteriol. 183:4823-4838(2001).
CC -!- FUNCTION: SbcCD cleaves DNA hairpin structures. These structures
CC can inhibit DNA replication and are intermediates in certain DNA
CC recombination reactions. The complex acts as a 3'->5' double
CC strand exonuclease that can open hairpins. It also has a 5'
CC single-strand endonuclease activity (By similarity).
CC -!- SUBUNIT: Heterodimer of sbcC and sbcD (By similarity).
CC -!- SIMILARITY: BELONGS TO THE SMC FAMILY. SBCC SUBFAMILY.
CC -----
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CC -----
CC DR EMBL; AE007771; AKK80682.1; -.
CC DR PIR; G97236; G97236.
CC DR InterPro; IPR003439; ABC transporter.
CC KW Hydrolyase; Nuclease; Exonuclease; Endonuclease; DNA replication;
CC DNA recombination; ATP-binding; Coiled coil; Complete proteome.
CC FT NP BIND 35 42 ATP (POTENTIAL).
CC FT DOMAIN 197 415 COILED COIL (POTENTIAL).
CC FT DOMAIN 446 1003 COILED COIL (POTENTIAL).
CC SQ SEQUENCE 1163 AA; 135507 MW; CE5F0BD2215D7A92 CRC64;

Query Match 78.9%; Score 30; DB 1; Length 1163;
Best Local Similarity 100.0%; Pred. No. 1.5e+02;
Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 INFEKL 8
Db 20 INFEKL 25

```

Search completed: January 30, 2004, 07:07:49
Job time : 14 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 30, 2004, 07:06:27 ; Search time 70 Seconds
(without alignments)
29.492 Million cell updates/sec

Title: SEQ10
Perfect score: 38
Sequence: 1 sinfexl 8

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database :

SPREMBL_23:*
1: sp_archaea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mhc:*
8: sp_organelle:*
9: sp_plant:*
10: sp_todent:*
11: sp_virus:*
12: sp_vertebrate:*
13: sp_unclassified:*
14: sp_virus:*
15: sp_bacteriap:*
16: sp_archaeap:*
17: sp_archaeap:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the total score being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	33	86.8	168	10 Q98RQ3	Q98RQ3 guillardi
2	33	86.8	315	10 O651S8	O651S8 phaseolus v
3	33	86.8	366	2 Q8VQ88	Q8VQ88 unclutered
4	32	84.2	260	16 Q9AA01	Q9AA01 caulobacter
5	32	84.2	299	16 Q92XR1	Q92XR1 rhizobium m
6	32	84.2	347	8 Q98RV6	Q98RV6 guillardi
7	32	84.2	443	17 Q8TRP5	Q8TRP5 mechanosarc
8	32	84.2	444	4 Q8IXU5	Q8IXU5 homo sapien
9	32	84.2	580	4 Q92565	Q92565 homo sapien
10	32	84.2	612	11 Q8BJT9	Q8BJT9 mus musculu
11	32	84.2	814	11 Q8CORS	Q8CORS mus musculu
12	32	84.2	814	11 Q8CQ09	Q8CQ09 mus musculu
13	32	84.2	965	2 Q9S4D1	Q9S4D1 staphylococ
14	32	84.2	965	2 Q8VVR0	Q8VVR0 staphylococ
15	32	84.2	1311	5 Q95NL8	Q95NL8 caenorhabdi
16	32	84.2	1347	5 Q95WR8	Q95WR8 caenorhabdi

17	32	84.2	1470	5 Q21218	Q21218 caenorhabdi
18	32	84.2	1573	5 Q9VWF3	Q9VWF3 drosophila
19	32	84.2	1573	5 Q95V18	Q95V18 drosophila
20	31	81.6	97	12 Q8BDP4	Q8BDP4 reindeer pa
21	31	81.6	103	5 Q8IB53	Q8IB53 plasmodium
22	31	81.6	211	5 P91370	P91370 caenorhabdi
23	31	81.6	287	11 Q8R1R1	Q8R1R1 mus musculu
24	31	81.6	307	17 Q95S82	Q95S82 pyrococcus
25	31	81.6	309	16 Q980R5	Q980R5 mycoplasma
26	31	81.6	364	10 Q9SKB4	Q9SKB4 arabidopsis
27	31	81.6	461	11 Q8BZK9	Q8BZK9 mus musculu
28	31	81.6	680	12 Q9Y7Q6	Q9Y7Q6 ateline her
29	31	81.6	834	11 Q8R3E5	Q8R3E5 mus musculu
30	31	81.6	876	11 Q8VCC8	Q8VCC8 mus musculu
31	31	81.6	881	4 Q8WVNO	Q8WVNO homo sapien
32	31	81.6	881	4 Q95634	Q95634 homo sapien
33	31	81.6	881	4 Q95398	Q95398 homo sapien
34	31	81.6	884	11 Q921C8	Q921C8 fatus norv
35	31	81.6	1089	5 Q81605	Q81605 plasmodium
36	31	81.6	1113	4 Q8TEA3	Q8TEA3 homo sapien
37	31	81.6	1138	11 Q8CHG7	Q8CHG7 mus saplen
38	31	81.6	1204	4 Q9UHV4	Q9UHV4 homo sapien
39	31	81.6	1391	4 Q8TEU6	Q8TEU6 homo sapien
40	31	81.6	1499	4 Q9Y4G8	Q9Y4G8 homo sapien
41	31	81.6	1509	4 Q96PC1	Q96PC1 homo sapien
42	31	81.6	1601	4 Q8TEU7	Q8TEU7 homo sapien
43	31	81.6	1601	4 Q8NT21	Q8NT21 homo sapien
44	31	81.6	1836	10 Q9LXR4	Q9LXR4 arabidopsis
45	31	81.6	1909	10 Q9LXR3	Q9LXR3 arabidopsis

ALIGNMENTS

RESULT 1					
ID	Q98RQ3	PRELIMINARY;	PRT;	168 AA.	
AC	Q98RQ3:				
DT	01-OCT-2001 (TREMBLrel. 18, Created)				
DT	01-OCT-2001 (TREMBLrel. 18, Last sequence update)				
DE	01-OCT-2001 (TREMBLrel. 18, Last annotation update)				
DE	Hypothetical 20.2 kDa protein orf168 from chromosome 1.				
GN	ORF168.				
OS	Guillardia theta (Cryptomonas phi).				
OC	Eukaryota; Cryptophyta; Cryptomonadaceae; Guillardia.				
OX	NCBI_TaxID=55529;				
RN	[1]				
RP	SEQUENCE FROM N.A.				
RX	MEDLINE=21223349; PubMed=11323671;				
RA	Douglas S. Zauner S., Fraunholz M., Beaton M., Penny S., Deng L.T.,				
RA	Wu X., Reich M., Cavalier-Smith T., Maier U.G.;				
RT	"The highly reduced genome of an enslaved algal nucleus."				
RL	Nature 410:1091-1096 (2001).				
DR	EMBL; AF165818; AAK39893.1; -				
KW	Hypothetical protein.				
SQ	SEQUENCE 168 AA; 20185 MW; 2874CBD53028A3D CRC64;				
Query Match					
Best Local Similarity 86.8%; Score 33; DB 10; Length 168;					
Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;					
QY	1 SINFEKL 8				
DB	108 NINFEKI 115				
RESULT 2					
ID	O651S8	PRELIMINARY;	PRT;	315 AA.	
AC	O651S8:				
DT	01-AUG-1998 (TREMBLrel. 07, Created)				
DT	01-AUG-1998 (TREMBLrel. 07, Last sequence update)				
DT	01-JUN-2002 (TREMBLrel. 21, Last annotation update)				

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DE 1-aminocyclopropane-1-carboxylic acid oxidase.
GN ACO1.
OS Phaeococcus vulgaris (kidney bean) (French bean).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eucosids I; Fabales; Fabaceae; Papilionoideae; Phaseoleae; Phaseolus.
OX NCBI_TaxID=3885;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Taylor; TISSUE=leaf;
RA Pidgeon C.M., Reid D.M., Facchini P.J.;
RT "Light induced changes in ethylene production in Phaeococcus vulgaris
RT cv. Taylor.";
RL Plant Physiol. 114:165-165(1997).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Taylor; TISSUE=leaf;
RA Pidgeon C.M., Facchini P.J., Reid D.M.;
RL Submitted (MAR-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF053354; AAC12934.1; -.
DR InterPro; IPR005123; 2OG-Fell_Oxy.
DR Pfam; PF03171; 2OG-Fell_Oxy; 1.
SQ SEQUENCE 315 AA; 36141 MW; 6898811DF88E9B81 CRC64;

Query Match
Best Local Similarity 86.8%; Score 33; DB 10; Length 315;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 IIINFEKL 8
DB 6 VINFEKL 12

RESULT 3
Q8VO88 PRELIMINARY; PRT; 366 AA.
AC Q8VO88;
DT 01-MAR-2002 (TREMBLrel. 20, Created)
DT 01-JUN-2002 (TREMBLrel. 21, Last sequence update)
DE Formyletrihydrofolate synthetase (Fragment).
OS uncultured environmental landfill bacterium.
OC Bacteria; environmental samples.
OX NCBI_TaxID=184235;
RN [1]
RP SEQUENCE FROM N.A.
RA Uz I., Lindner A.S., Rasche M.E., Townsend T.G., Ogram A.V.;
RT "Microbial Characterization of Landfill Cover Soils of Different
RT Ages.";
RL Submitted (DEC-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF459462; AAL67505.1; -.
DR InterPro; IPR000559; Fmtehyd_synth.
DR Pfam; PF01268; FTHFS; 1.
FT NON TER 1 1
FT NON TER 366 366
SQ SEQUENCE 366 AA; 39816 MW; EB4685D501E97026 CRC64;

Query Match
Best Local Similarity 86.8%; Score 33; DB 2; Length 366;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 IIINFEKL 8
DB 346 IVNFEKL 352

RESULT 4
Q9AA01 PRELIMINARY; PRT; 260 AA.
AC Q9AA01;
DT 01-JUN-2001 (TREMBLrel. 17, Created)
DT 01-JUN-2001 (TREMBLrel. 17, Last sequence update)
DT 01-JUN-2002 (TREMBLrel. 21, Last annotation update)

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DE Hypothetical protein CC0810.
GN CC0810.
OS Caulobacter crescentus.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Caulobacteriales;
OC Caulobacteriaceae; Caulobacter.
OX NCBI_TaxID=155892;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC 19089 / CB15;
RX MEDLINE=21173698; PubMed=11259647;
RA Nierman W.C., Feldblyum T.V., Laud M.T., Paulsen I.T., Nelson K.E.,
RA Eisen J., Heidelberg J.F., Alley M.R.K., Ohta N., Maddock J.R.,
RA Potocka I., Nelson W.C., Newton A., Stephens C., Phadke N.D., Ely B.,
RA DeBoy R.T., Dodson R.J., Durkin A.S., Gwinn M.L., Haft D.H.,
RA Kolonay J.F., Smit J., Craven M.B., Khouri H., Shetty J., Berry K.,
RA Ueberbach T., Tran K., Wolf A., Vamathavan J., Ermlaeva M., White O.,
RA Salzberg S.L., Venter J.C., Shapiro L., Fraser C.M.;
RT "Complete genome sequence of Caulobacter crescentus.";
RL Proc. Natl. Acad. Sci. U.S.A. 98:4136-4141(2001).
DR EMBL; AE005757; AAK22795.1; -.
DR TIGR; CC0810; -.
DR InterPro; IPR005269; Cons_hypoth730.
DR Pfam; PF03641; Lysine_decarbox; 1.
DR TIGRFAMs; TIGR00730; TIGR00730; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 260 AA; 28789 MW; E4CF8523B12EF1B CRC64;

Query Match
Best Local Similarity 84.2%; Score 32; DB 16; Length 260;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 IIINFEKL 8
DB 215 SIINFEAL 222

RESULT 5
Q9ZKR1 PRELIMINARY; PRT; 299 AA.
AC Q9ZKR1;
DT 01-DEC-2001 (TREMBLrel. 19, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DT 01-JUN-2002 (TREMBLrel. 21, Last annotation update)
DE Hypothetical protein R00434.
GN R00434 OR SMC01731.
OS Rhizobium meliloti (Sinorhizobium meliloti).
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Rhizobiaceae; Sinorhizobium.
OX NCBI_TaxID=382;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=1021;
RX MEDLINE=21396507; PubMed=11481430;
RA Capela D., Barloy-Hubler F., Gouzy J., Bothe G., Ampe F., Batut J.,
RA Boistard P., Becker A., Boutry M., Cadieu E., Dreaou S., Gloux S.,
RA Godie T., Goffeau A., Kahn D., Kiss E., Lelaune V., Masny D.,
RA Pohl T., Portetelle D., Puehler A., Purnelle B., Ramberger U.,
RA Renard C., Thebaud P., Vandembol M., Weidner S., Galibert F.;
RT "Analysis of the chromosome sequence of the legume symbiont
RT Sinorhizobium meliloti strain 1021.";
RL Proc. Natl. Acad. Sci. U.S.A. 98:9877-9882(2001).
DR EMBL; AL591783; CAC41871.1; -.
DR InterPro; IPR005269; Cons_hypoth730.
DR Pfam; PF03641; Lysine_decarbox; 1.
KW Hypothetical protein; Complete proteome.
SQ SEQUENCE 299 AA; 33440 MW; 9BD61375B91F195 CRC64;

Query Match
Best Local Similarity 84.2%; Score 32; DB 16; Length 299;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 IIINFEKL 8

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Db 246 STINEAL 253

RESULT 6

ID Q98RV6 PRELIMINARY; PRT; 347 AA.

AC Q98RV6;

DT 01-OCT-2001 (TRENBLrel. 18, Created)

DT 01-OCT-2001 (TRENBLrel. 18, Last sequence update)

DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)

DE Cyclin B.

GN CYCB.

OS Guillardia theta (Cryptomonas phi).

OC Nucleomorph.

OC Eukaryota; Cryptophyta; Cryptomonadaceae; Guillardia.

OX NCBI_TaxID=55529;

RN [1]

RP SEQUENCE FROM N.A.

RX MEDLINE=2123349; PubMed=1123671;

RA Douglas S., Zauner S., Fraunholz M., Beaton M., Penny S., Deng L.T.,

Wu X., Reith M., Cavalier-Smith T., Maiter U.G.;

RT "The highly reduced genome of an enslaved algal nucleus."

RL Nature 410:1091-1096(2001).

CC -1. SIMILARITY: BELONGS TO THE CYCLIN FAMILY.

DR EMBL; AF165818; AAK39844.1; -.

DR InterPro; IPR006670; Cyclin.

DR InterPro; IPR006671; Cyclin_N.

DR Pfam; PF00134; Cyclin; 1.

DR SMART; SM00385; CYCLIN; 2.

DR PROSITE; PS00292; CYCLINS; 1.

DR PROSITE; Cell division; Cyclin.

DR CELL cycle; Cell division; Cyclin.

SQ SEQUENCE 347 AA; 41616 MW; CC579909C403A04D CRC64;

QY

Db

1 STINEFKL 8

86 NVLNFEKL 93

RESULT 7

O8TRP5

ID O8TRP5 PRELIMINARY; PRT; 443 AA.

AC O8TRP5;

DT 01-UN-2002 (TRENBLrel. 21, Created)

DT 01-UN-2002 (TRENBLrel. 21, Last sequence update)

DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)

DE Iron ABC transporter, solute-binding protein.

GN MA1130.

OS Methanosarcina acetivorans.

OC Archaea; Euryarchaeota; Methanococci; Methanosarcinales;

OC Methanosarcinaceae; Methanosarcina.

OX NCBI_TaxID=2214;

RN [1]

RP SEQUENCE FROM N.A.

RX STRAIN=C2A / ATCC 35395 / DSM 2834;

RA MEDLINE=21929760; PubMed=11932238;

RA Galagan J.E., Nussbaum C., Roy A., Endrizzi M.G., MacDonald P.,

RA Fitzhugh W., Calvo S., Engels R., Smitrov S., Anoor D., Brown A.,

RA Allen N., Naylor J., Stange-Thomann N., Dearellano K., Johnson R.,

RA Linton L., McKean P., McKernan K., Talamas J., Turrell A., Ye W.,

RA Zimmer A., Barber R.D., Cam I., Graham D.E., Grahame D.A., Guss A.M.,

RA Hedderich R., Ingram-Smith C., Kuetner H.C., Krzycki J.A.,

RA Leigh J.A., Li W., Liu J., Mukhopadhyay B., Reeve J.N., Smith K.,

RA Springer T.A., Umayam L.A., White O., White R.H., de Macario E.C.,

RA Ferry J.G., Jarrell K.F., Jung H., Macario A.J.L., Paulsen I.,

RA Pritchett M., Sowers K.R., Swanson R.V., Zinder S.H., Lander E.,

RA Metcalf W.W., Birren B.,

RT "The genome of Methanosarcina acetivorans reveals extensive metabolic

RT and physiological diversity."

RL Genome Res. 12:532-542(2002).

DR EMBL; AE010779; AAK04551.1; -.

DR InterPro; IPR002491; Peripla_BP.

DR Pfam; PF01497; Peripla_BP_2; 1.

KW Complete proteome.

SQ SEQUENCE 443 AA; 49537 MW; 0D3F2198B20E86C9 CRC64;

QY

Db

1 STINEFKL 8

166 TIINYEKL 173

RESULT 8

O8IXU5

ID O8IXU5 PRELIMINARY; PRT; 444 AA.

AC O8IXU5;

DT 01-MAR-2003 (TRENBLrel. 23, Created)

DT 01-MAR-2003 (TRENBLrel. 23, Last sequence update)

DT 01-MAR-2003 (TRENBLrel. 23, Last annotation update)

DE Similar to guanine nucleotide exchange factor for Rap1,

DE M-Ras-regulated GEF.

DE Homo sapiens (Human).

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

OX NCBI_TaxID=9606;

RN [1]

RP SEQUENCE FROM N.A.

RX TISSUE=Brain;

RA Strausberg R.;

RL Submitted (NOV-2002) to the EMBL/GenBank/DBJ databases.

DR EMBL; BC039203; AAH39203.1; -.

SQ SEQUENCE 444 AA; 52051 MW; ECEBDEP820DDFF13B CRC64;

QY

Db

1 STINEFKL 8

381 NVLNFEKL 388

RESULT 9

O92565

ID O92565 PRELIMINARY; PRT; 580 AA.

AC O92565;

DT 01-FEB-1997 (TRENBLrel. 02, Created)

DT 01-FEB-1997 (TRENBLrel. 02, Last sequence update)

DT 01-OCT-2002 (TRENBLrel. 22, Last annotation update)

DE Hypothetical protein KIAA0277.

GN KIAA0277.

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

OX NCBI_TaxID=9606;

RN [1]

RP SEQUENCE FROM N.A.

RX TISSUE=Brain;

RA MEDLINE=97191544; PubMed=9039502;

RA Nagase T., Seki N., Ishikawa K., Ohira M., Kawarabayashi Y., Ohara O.,

RA Tanaka A., Kotani H., Miyajima N., Nomura N.;

RT "Prediction of the coding sequences of unidentified human genes. VI.

RT the coding sequences of 80 new genes (KIAA0201-KIAA0280) deduced by

RT analysis of cDNA clones from cell line KG-1 and brain."

RL DNA Res. 3:321-329(1996).

DR EMBL; D87467; BA013406.1; -.

DR InterPro; IPR000651; RasGEFN.

DR InterPro; IPR001895; RasGRF_CDC25.

DR Pfam; PF00617; RasGEFN; 1.

DR Pfam; PF00618; RasGEFN; 1.

DR SMART; SM00147; RASGEF; 1.
 DR SMART; SM00229; RASGEF; 1.
 KW Hypothetical protein.
 SQ SEQUENCE 580 AA; 67733 MW; 732PB7A1IDFPA1C CRC64;

Query Match
 Best Local Similarity 84.2%; Score 32; DB 4; Length 580;
 Pred. No. 2.1e+02;
 Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINPEKL 8
 ::::|
 DB 517 NLVNFEXL 524

RESULT 10
 Q8BJ9 PRELIMINARY; PRT; 612 AA.
 ID Q8BJ9
 AC Q8BJ9;
 DT 01-MAR-2003 (TrEMBLrel. 23, Created)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE Hypothetical guanine-nucleotide dissociation stimulators CDC25
 DE family/guanine nucleotide exchange factor for Ras-like GTPases;
 DE N-terminal motif containing protein.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Body;
 RX MEDLINE=22354683; PubMed=12466851;
 RA The FANTOM Consortium;
 RT the RIKEN Genome Exploration Research Group Phase I & II Team;
 RT "Analysis of the mouse transcriptome based on functional annotation of
 60,770 full-length cDNAs."
 RL Nature 420:563-573 (2002).
 DR EMBL; AK083591; BAC38963.1; -.
 KW Hypothetical protein.
 SQ SEQUENCE 612 AA; 70913 MW; DBD552DDFD392B6 CRC64;

Query Match
 Best Local Similarity 84.2%; Score 32; DB 11; Length 612;
 Pred. No. 2.2e+02;
 Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINPEKL 8
 ::::|
 DB 548 NLVNFEXL 555

RESULT 11
 Q8C0R5 PRELIMINARY; PRT; 814 AA.
 ID Q8C0R5
 AC Q8C0R5;
 DT 01-MAR-2003 (TrEMBLrel. 23, Created)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE Hypothetical guanine-nucleotide dissociation stimulators CDC25
 DE family/guanine nucleotide exchange factor for Ras-like GTPases;
 DE N-terminal motif containing protein.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Testis;
 RX MEDLINE=22354683; PubMed=12466851;
 RA The FANTOM Consortium;
 RT the RIKEN Genome Exploration Research Group Phase I & II Team;
 RT "Analysis of the mouse transcriptome based on functional annotation of
 60,770 full-length cDNAs."
 RL Nature 420:563-573 (2002).

DR EMBL; AK029995; BAC26723.1; -.
 KW Hypothetical protein.
 SQ SEQUENCE 814 AA; 93681 MW; A6490A72AA3E89B CRC64;

Query Match
 Best Local Similarity 84.2%; Score 32; DB 11; Length 814;
 Pred. No. 2.9e+02;
 Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINPEKL 8
 ::::|
 DB 750 NLVNFEXL 757

RESULT 12
 Q8C0Q9 PRELIMINARY; PRT; 814 AA.
 ID Q8C0Q9
 AC Q8C0Q9;
 DT 01-MAR-2003 (TrEMBLrel. 23, Created)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
 DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
 DE Hypothetical guanine-nucleotide dissociation stimulators CDC25
 DE family/guanine nucleotide exchange factor for Ras-like GTPases;
 DE N-terminal motif containing protein.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57BL/6J; TISSUE=Testis;
 RX MEDLINE=22354683; PubMed=12466851;
 RA The FANTOM Consortium;
 RT the RIKEN Genome Exploration Research Group Phase I & II Team;
 RT "Analysis of the mouse transcriptome based on functional annotation of
 60,770 full-length cDNAs."
 RL Nature 420:563-573 (2002).
 DR EMBL; AK030016; BAC26736.1; -.
 KW Hypothetical protein.
 SQ SEQUENCE 814 AA; 93754 MW; AA04EBC74CE13735 CRC64;

Query Match
 Best Local Similarity 84.2%; Score 32; DB 11; Length 814;
 Pred. No. 2.9e+02;
 Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINPEKL 8
 ::::|
 DB 750 NLVNFEXL 757

RESULT 13
 Q9S4D1 PRELIMINARY; PRT; 965 AA.
 ID Q9S4D1
 AC Q9S4D1;
 DT 01-MAY-2000 (TrEMBLrel. 13, Created)
 DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
 DT 01-DEC-2001 (TrEMBLrel. 19, Last annotation update)
 DE Lanthibiotic modifying enzyme.
 OS Staphylococcus aureus.
 OC Bacteria; Firmicutes; Bacillales; Staphylococcus.
 OX NCBI_TaxID=1280;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C57;
 RX MEDLINE=99346225; PubMed=104117203;
 RA Navaracna M.A., Sahl H.G., Tagg J.R.,
 RT "Identification of genes encoding two-component lanthibiotic production
 in Staphylococcus aureus C55 and other phage group II S. aureus
 strains and demonstration of an association with the exfoliative toxin
 B gene."
 RT Infect. Immun. 67:4268-4271 (1999).
 DR EMBL; AF147744; AAD47013.1; -.
 SQ SEQUENCE 965 AA; 111570 MW; BC1EADABDAC4F346 CRC64;

Query Match 84.2%; Score 32; DB 2; Length 965;
 Best Local Similarity 71.4%; Pred. No. 3.4e+02;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEX 7
 DB 202 SVNFEX 208

RESULT 14

OSVVR0 PRELIMINARY; PRT; 965 AA.
 AC OSVVR0;
 DT 01-MAR-2002 (TREMBlrel. 20, Created)
 DT 01-MAR-2002 (TREMBlrel. 20, Last sequence update)
 DT 01-MAR-2002 (TREMBlrel. 20, Last annotation update)
 DE ORF2.
 OS Staphylococcus aureus.
 OG Plasmid ETB plasmid.
 OC Bacteria; Firmicutes; Bacillales; Staphylococcus.
 OX NCBI_Taxid=1280;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=TY4;
 RX MEDLINE=21562640; PubMed=11705958;
 RA Yamaguchi T., Hayashi T., Takami H., Ohnishi M., Murata T.,
 RA Nakayama K., Asakawa K., Ohara M., Komatsuza H., Sugai M.,
 RT "Complete Nucleotide Sequence of a Staphylococcus aureus Exfoliative
 RT Toxin B Plasmid and Identification of a Novel ADP-Ribosyltransferase,
 RT EDIN-C.";
 RL Infect. Immun. 69:7760-7771(2001).
 DR EMBL; AF003088; BAB78440.1; -.
 KW Plasmid.
 SQ SEQUENCE 965 AA; 111561 MW; C08E9FD8DAC5F82B CRC64;

Query Match 84.2%; Score 32; DB 2; Length 965;
 Best Local Similarity 71.4%; Pred. No. 3.4e+02;
 Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEX 7
 DB 202 SVNFEX 208

RESULT 15
 Q95N18 PRELIMINARY; PRT; 1311 AA.
 AC Q95N18;
 DT 01-DEC-2001 (TREMBlrel. 19, Created)
 DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
 DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
 DE T14G10.2b protein (PKF isoform B).
 GN T14G10.2 OR T14G10.2B.
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
 OC Rhabditidae; Peloderinae; Caenorhabditis.
 OX NCBI_Taxid=6239;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Wild A.;
 RL Submitted (JAN-1996) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=99069613; PubMed=9851916;
 RA none;
 RT "Genome sequence of the nematode C.elegans: A platform for
 RT investigating biology.";
 RL Science 283:2012-2018(1998).
 RN [3]
 RP SEQUENCE FROM N.A.
 RA Vermeulen M., van Berkel W., Jansen G., de Rooij J., Plasterk R.H.,
 RA Bos J.L., Zwartkruis F.J.T.;
 RT "Characterization of pxf, the C. elegans homolog of human PDZ-GFPs.";

RL Submitted (SEP-2000) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.

RA Wild A.;
 RL Submitted (FEB-1996) to the EMBL/GenBank/DBJ databases.
 CC -1 SIMILARITY: CONTAINS 1 PDZ/DHR DOMAIN.
 DR EMBL; 268880; CAC42342.1; -.
 DR EMBL; 269664; CAC42342.1; JOINED.
 DR EMBL; AF308448; AAL09434.1; -.
 DR EMBL; 269664; CAC42313.1; -.
 DR EMBL; 268880; CAC42313.1; JOINED.
 DR Wormpep; T14G10.2b; CE28081.
 DR InterPro; IPR000595; CNMP_binding.
 DR InterPro; IPR001478; PDZ.
 DR InterPro; IPR000651; RASGEFN.
 DR InterPro; IPR001895; RASGEF_CDC25.
 DR InterPro; IPR00159; RA_domain.
 DR Pfam; PF00027; CNMP_binding; 2.
 DR Pfam; PF00595; PDZ; 1.
 DR Pfam; PF00789; RA; 1.
 DR Pfam; PF00617; RASGEF; 1.
 DR SMART; SM00100; CNMP; 1.
 DR SMART; SM00228; PDZ; 1.
 DR SMART; SM00314; RA; 1.
 DR SMART; SM00147; RASGEF; 1.
 DR SMART; SM00229; RASGEFN; 1.
 DR PROSITE; PS50042; CNMP_BINDING_3; 1.
 DR PROSITE; PS50106; PDZ_1.
 SQ SEQUENCE 1311 AA; 147003 MW; A4870F07DC201F97 CRC64;

Query Match 84.2%; Score 32; DB 5; Length 1311;
 Best Local Similarity 85.7%; Pred. No. 4.5e+02;
 Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 IINFEXL 8
 DB 1068 LINFEXL 1074

Search completed: January 30, 2004, 07:09:11
 Job time : 71 secs

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OM protein - protein search, using sw model

Run on: January 30, 2004, 07:15:53 ; Search time 71 Seconds

(without alignments)
17,885 Million cell updates/sec

Title: SEQ10

Perfect score: 38

Sequence: 1 sinifexl 8

Scoring table: BLOSUM62

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Searched: 1107863 seqs, 156726573 residues

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Post-processing: Minimum Match 100%

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Listing first 250 summaries

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21: /SIDSI/gcgdata/geneseq/geneseqp-emb1/AA2000.DAT.*
22: /SIDSI/gcgdata/geneseq/geneseqp-emb1/AA2001.DAT.*
23: /SIDSI/gcgdata/geneseq/geneseqp-emb1/AA2002.DAT.*
24: /SIDSI/gcgdata/geneseq/geneseqp-emb1/AA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	38	100.0	8	15	AAK57996
2	38	100.0	8	16	AAK83938
3	38	100.0	8	17	AAK89157
4	38	100.0	8	18	AAK19955
5	38	100.0	8	18	AAK14087
6	38	100.0	8	18	AAK04642
7	38	100.0	8	19	AAK80296
8	38	100.0	8	19	AAK70375
9	38	100.0	8	19	AAK68308

10	38	100.0	8	19	AAK68365	MHC binding peptid
11	38	100.0	8	19	AAK60700	Ovalbumin peptide
12	38	100.0	8	19	AAK54265	Ovalbumin peptide
13	38	100.0	8	19	AAK52838	Ovalbumin artifact
14	38	100.0	8	20	AAK42307	Ovalbumin-derived
15	38	100.0	8	20	AAK16871	OVA peptide sequen
16	38	100.0	8	20	AAK03780	Ovalbumin peptide
17	38	100.0	8	20	AAK99480	Ovalbumin-derived
18	38	100.0	8	20	AAK67585	Ovalbumin-derived
19	38	100.0	8	21	AAK13763	T-cell activation
20	38	100.0	8	21	AAK29465	Peptide fragment f
21	38	100.0	8	21	AAK26484	Negative control p
22	38	100.0	8	21	AAK3785	Tumour associated
23	38	100.0	8	21	AAK68311	OVA-derived peptid
24	38	100.0	8	21	AAK59401	Altered MHC determ
25	38	100.0	8	21	AAK52564	Ovalbumin protein
26	38	100.0	8	21	AAK52965	Marine ovalbumin M
27	38	100.0	8	22	AAK77871	Altered MHC determ
28	38	100.0	8	22	AAK52562	Ovalbumin 257-264
29	38	100.0	8	22	AAK13119	Cytotoxic T-cell e
30	38	100.0	8	22	AAK12145	Ovalbumin (OVA)-de
31	38	100.0	8	22	AAK09514	Marine ovalbumin (
32	38	100.0	8	22	AAK63855	Human ovalbumin pe
33	38	100.0	8	22	AAK66422	Amino acid sequenc
34	38	100.0	8	22	AAK05398	Chicken ovalbumin
35	38	100.0	8	22	AAK06033	Peptide released f
36	38	100.0	8	22	AAK84316	Chicken ovalbumin
37	38	100.0	8	22	AAK99354	Peptide used to pr
38	38	100.0	8	22	AAK82176	Ovalbumin cytotoxi
39	38	100.0	8	22	AAK81122	Immunodominant CTL
40	38	100.0	8	22	AAK82065	Chicken ovalbumin
41	38	100.0	8	22	AAK92374	Ovalbumin-derived
42	38	100.0	8	22	AAK82374	Miscellaneous pept
43	38	100.0	8	22	AAK8950	Ovalbumin MHC clas
44	38	100.0	8	23	AAK28959	Chicken ovalbumin
45	38	100.0	8	23	AAK26368	Ovalbumin CTL epit
46	38	100.0	8	23	AAK79933	Ovalbumin T-cell e
47	38	100.0	8	23	AAK93028	Mouse class I MHC
48	38	100.0	8	23	AAK25400	Ovalbumin peptide
49	38	100.0	8	23	AAK31661	Chicken ovalbumin
50	38	100.0	8	23	AAK31967	Chicken ovalbumin
51	38	100.0	8	23	AAK99718	Mouse MHC class I K
52	38	100.0	8	23	AAK08108	Chicken ovalbumin
53	38	100.0	8	23	AAK81273	Chicken OVA 257-26
54	38	100.0	8	23	AAK22531	Ovalbumin peptide
55	38	100.0	8	23	AAK76050	Ovalbumin, H-2kb r
56	38	100.0	8	23	AAK19945	Cytotoxic T-cell e
57	38	100.0	8	23	AAK09907	Ovalbumin peptide
58	38	100.0	8	23	AAK76942	OVA peptide (257-2
59	38	100.0	8	23	AAK76802	MHC class I-restri
60	38	100.0	8	23	AAK1866	OVA peptide fragme
61	38	100.0	8	23	AAK11239	Ovalbumin derived c
62	38	100.0	8	23	AAK09820	Immunodominant Kb-
63	38	100.0	8	23	AAK33215	Ovalbumin-derived
64	38	100.0	8	23	AAK13436	Hen egg ovalbumin
65	38	100.0	8	24	AAK75056	Chicken ovalbumin
66	38	100.0	8	24	AAK07743	Ovalbumin antigeni
67	38	100.0	8	24	AAK08619	Chicken ovalbumin
68	38	100.0	8	24	AAK7401	Ovalbumin (OVA) re
69	38	100.0	8	24	AAK58359	Synthetic bmer pep
70	38	100.0	8	24	ABG73081	Ovalbumin-derived
71	38	100.0	8	24	ABP6760	MHC class I peptid
72	38	100.0	8	24	ABP60027	Ovalbumin derived
73	38	100.0	8	24	ABU11029	Ovalbumin antigeni
74	38	100.0	9	24	AAK4323	Ovalbumin immunodo
75	38	100.0	9	24	ABP57402	Peptide used to pr
76	38	100.0	10	18	AAK04643	Synthetic bmer pep
77	38	100.0	10	18	AAK04644	Ovalbumin-derived
78	38	100.0	10	23	AAK09821	Ovalbumin-derived
79	38	100.0	10	23	AAK09825	Modified ovalbumin
80	38	100.0	12	23	AAK14122	Modified ovalbumin
81	38	100.0	12	23	AAK09822	OVA protein derive
82	38	100.0	23	23	AAK09826	Modified ovalbumin

83	38	100.0	12	23	AAU09827	Modified ovalbumin
84	38	100.0	14	23	ABB76049	Peptide insert in
85	38	100.0	14	23	AAU09823	Modified ovalbumin
86	38	100.0	14	23	AAU09828	Modified ovalbumin
87	38	100.0	15	23	AAU09824	Modified ovalbumin
88	38	100.0	16	24	ABP57403	Synthetic lemer pe
89	38	100.0	19	18	AAW19957	B1P-binding domain
90	38	100.0	19	18	AAW19956	OVA-B1P-binding do
91	38	100.0	19	23	AAE13446	Chicken MHC class
92	38	100.0	19	23	AAE13447	Chicken MHC class
93	38	100.0	19	24	ABP57404	Synthetic lemer pe
94	38	100.0	24	14	AAE32294	Synthetic peptide
95	38	100.0	24	14	AAE41450	Antigenic peptide
96	38	100.0	24	18	AAW04645	Ovalbumin-derived
97	38	100.0	24	12	AAW65107	Ovalbumin based pe
98	38	100.0	24	22	ABP57439	Myelin basic prote
99	38	100.0	24	23	ABG31664	Ovalbumin (OVA) pe
100	38	100.0	26	24	ABP57405	Synthetic 26mer pe
101	38	100.0	26	24	ABP57406	Synthetic 26mer* p
102	38	100.0	30	23	AAE13448	Chicken MHC class
103	38	100.0	31	24	ABP57407	Synthetic 31mer pe
104	38	100.0	35	18	AAW04646	Ovalbumin-derived
105	38	100.0	36	24	AAO26741	Chicken ovalbumin
106	38	100.0	43	22	AAE84325	Amino acid sequenc
107	38	100.0	47	22	AAE84321	Amino acid sequenc
108	38	100.0	48	22	AAE84322	Amino acid sequenc
109	38	100.0	49	22	AAE84953	T5-DICE ovalbumin
110	38	100.0	57	22	AAE84954	DICE-I ovalbumin M
111	38	100.0	100	23	AAE13458	Chicken ovalbumin
112	38	100.0	100	23	AAE13460	Chicken ovalbumin
113	38	100.0	103	23	AAE13459	Chicken ovalbumin
114	38	100.0	103	23	AAE13461	Chicken ovalbumin
115	38	100.0	106	17	AAE89966	Polytope sequence.
116	38	100.0	108	23	AAE13462	Chicken ovalbumin
117	38	100.0	111	23	AAE13463	Chicken ovalbumin
118	38	100.0	132	21	AAE52575	Amino acid sequenc
119	38	100.0	386	23	AAE13435	Chicken ovalbumin
120	38	100.0	409	22	AAE13435	Amino acid sequenc
121	38	100.0	479	22	AAE13112	Human HER300-rGM-C
122	38	100.0	541	23	AAU99725	Yeast/mouse SS-OVA
123	38	100.0	564	22	AAE13110	Human HER500 fusio
124	38	100.0	697	22	AAE13111	Human HER500-rGM-C
125	38	100.0	948	22	AAE31611	Amino acid sequenc

ALIGNMENTS

RESULT 1
AAR57996
ID AAR57996 standard; Protein; 8 AA.
XX
AC AAR57996;
XX
DT 25-MAR-2003 (updated)
DT 30-MAR-1995 (first entry)
XX
DE Ova257-264.
XX
XX Ova; ovalbumin; cytosol; cytolitic immune response;
KW vaccinia virus; promoter.
OS Synthetic.
XX
XX WO9417816-A1.
XX
XX 18-AUG-1994.
XX
XX 27-JAN-1994; 94WO-US01183.
XX
XX 10-FEB-1993; 93US-0016066.
XX
XX (DAND) DANA FARBER CANCER INST INC.

PA (HARD) HARVARD COLLEGE.
XX Goldberg AL, Rock KL;
PI
XX WPI; 1994-279383/34.
DR
XX
XX
XX Method for blocking cytolitic immune responses - useful for
PT treatment of autoimmune diseases and preventing organ and graft
PT rejection
XX
XX
XX Example 2; Page 44; 89pp; English.
XX
XX Ova257-264 was constructed by inserting a synthetic oligonucleotide
CC (AAO6742) behind the vaccinia virus p7.5 early/late promoter in psc11
CC which was modified such that the restriction sites SalI and NotI
CC were substituted for the SmaI site. The oligonucleotide consists of
CC a SalI site, Kozak's consensus sequence for efficient translation,
CC an initiation codon, nucleotides encoding the peptide given in
CC AAR57996, two stop codons, and a NotI site.
CC (Updated on 25-MAR-2003 to correct PN field.)
XX
SQ Sequence 8 AA;
Query Match 100.0%; Score 38; DB 15; Length 8;
Best local similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 STINPEKL 8
Db 1 STINPEKL 8
RESULT 2
AAR83938
ID AAR83938 standard; peptide; 8 AA.
XX
AC AAR83938;
XX
DT 05-JUN-1996 (first entry)
XX
DE MHC class I restricted antigenic peptide #8.
XX
XX MHC class I; antigen; MAGE; melanoma; breast cancer; bladder cancer;
KW Titermax; cytotoxic T-lymphocyte; tumour; pathogenic disease; bacteria;
KW parasite; human; animal.
OS Synthetic.
XX
XX WO9528958-A1.
XX
XX 02-NOV-1995.
XX
XX 21-APR-1995; 95WO-US04975.
XX
XX 22-APR-1994; 94US-0233496.
XX
XX (SLOK) SLOAN KETTERING INST CANCER RES.
XX
XX Dyall R, Nikolic-Zugic J;
PI
XX WPI; 1995-382848/49.
XX
XX
XX Cytotoxic T-cell induction by MHC class I-restricted peptide in
PT adjuvant - useful for treating tumours and bacterial or parasitic
PT pathogenic diseases
XX
XX
XX Claim 11; Page 38; 50pp; English.
XX
XX The sequences given in AAR83931-49 are MHC class I restricted 8-12
CC amino acid antigenic peptides. This peptide represents ovalbumin
CC residues 257-264. These peptides may be administered to a subject
CC in combination with a suitable adjuvant, pref. Titermax (RTM), to
CC induce cytotoxic T-lymphocytes. This method may be used in the

CC treatment of a tumour or a pathogenic disease, esp. diseases of
 CC bacterial or parasitic origin, in humans and animals, e.g monkeys,
 CC dogs, cows, horses, etc.
 XX

SO Sequence 8 AA;

Query Match 100.0%; Score 38; DB 16; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEKL 8
 |||||
 Db 1 SIINFEKL 8

RESULT 3

AA89157
 ID AA89157 standard; peptide; 8 AA.

AA89157;

25-MAR-2003 (updated)
 03-SEP-1996 (first entry)

Peptide Ova8 used in an MHC stripping/reloading method.

Antigen; major histocompatibility complex; cell surface; stimulation;
 cytotoxic T lymphocyte; CTL; endogenous; exogenous; peptide; spleen;
 allelic restriction; peripheral blood lymphocytes; ganglion; placenta;
 native form; infection; tumour; autoimmune disease.

Synthetic.

MO9601891-A1.

25-JAN-1996.

06-JUL-1995; 95MO-FR00907.

07-JUL-1994; 94FR-0008427.

(INRM) INSERM INST NAT SANTE & RECH MEDICALE.

(INSP) INST PASTEUR.

Langlade Demoyen P, Kourilsky P, Abastado J;

WPI; 1996-097621/10.

Cell population with high surface density of exogenous peptide bound
 to MHC molecules - prepd. by stripping endogenous peptide and
 reloading exogenous peptide(s), useful for stimulating cytotoxic
 lymphocytes in cases of infection, tumour and auto-immune disease

Disclosure; Page 9; 37pp; French.

Peptides AAR89154-73 are examples of exogenous peptides which are
 "loaded" onto the antigen-presenting major histocompatibility complex
 (MHC) on the cell surface in a novel method of stimulating cytotoxic T
 lymphocytes. The method involves treating cells, e.g. at a pH 5 or below
 or pH 9 or above, to remove the endogenous peptides from the MHC,
 followed by recharging the complexes with specified exogenous peptides
 having the same allelic restriction as the MHC. The cells are pref.
 peripheral blood lymphocytes, spleen, ganglion or placental cells which
 are able to present the exogenous antigen in a native form. Recharged
 cells contain exogenous peptides at a higher density than native cells
 and are used to stimulate specific cytotoxic T lymphocytes in response to
 infections, tumours or autoimmune diseases. This peptide is derived from
 ovalbumin and belongs to the allelic restriction Kb.
 (Updated on 25-MAR-2003 to correct PI field.)

Sequence 8 AA;

Query Match 100.0%; Score 38; DB 17; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEKL 8
 |||||
 Db 1 SIINFEKL 8

RESULT 4

AAW19955
 ID AAW19955 standard; Peptide; 8 AA.

AAW19955;

10-NOV-1997 (first entry)

Chicken OVA-peptide.

Vaccine; immunotherapy; heat shock protein; OVA; cancer;
 infectious disease.

Gallus sp.

MO9706821-A1.

27-FEB-1997.

16-AUG-1996; 96MO-US13363.

18-AUG-1995; 95US-0002490.

18-AUG-1995; 95US-0002479.

(SLOK) SLOAN KETTERING INST CANCER RES.

Hartl FU, Hoe MH, Houghton A, Mayhew M, Rothman JE;

Takeuchi Y;

WPI; 1997-165035/15.

Compsn. for inducing immune response contg. antigen and heat shock
 protein - also new hybrid peptide and related nucleic acid, for
 treatment of infectious diseases and tumours

Example 1; Page 17; 58pp; English.

Chicken OVA-peptide (AAW19955) is used in novel hybrid peptides,
 OVA-BiP (AAW19956) and BiP-OVA (AAW19957) with heat shock protein (HSP)
 BiP binding domain (see also AAW19951). The hybrid protein is
 combined in vitro with a HSP, such as hsp70, to form a complex
 that, when administered to a subject, induces an immune response.
 Vaccine compositions were prepd. by combining recombinant mouse
 hsp70, recombinant human hsp40 and Ova-peptide. Combinations of
 CC antigen with hsp70 or a mixture of hsp70 and hsp40 were effective
 to produce a cytotoxic T lymphocyte response.

Sequence 8 AA;

Query Match 100.0%; Score 38; DB 18; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEKL 8
 |||||
 Db 1 SIINFEKL 8

RESULT 5

AAW14087
 ID AAW14087 standard; peptide; 8 AA.

AAW14087;

20-OCT-1997 (first entry)

```

XX MHC class I molecule binding OVA peptide.
DE
XX
XX Major histocompatibility complex; MHC; target; binding; tumour;
KM cancer; neoplasia; LSTRA; EL-4; identification; detection; screening;
KM tissue typing; Bcr-abl.
XX
XX Mus sp.
OS
XX WO9641188-A1.
PN
XX 19-DEC-1996.
PD
XX 07-JUN-1996; 96WO-US09680.
PF
XX 07-JUN-1995; 95US-0485610.
PR
XX (UNITM ) UNIV WASHINGTON.
PA
XX Cheever MA, Chen W;
PI
XX WPI; 1997-108657/10.
DR
XX
XX Identifying major histocompatibility complex class I binding mols. -
PT using peptide(s) having a core of 7-14 amino acids with extra amino
PT acids and a reporter gp. at the N- or C-terminus, useful for tissue
PT typing
XX
XX Example 2; Page 19; 41pp; English.
PS
XX AAW44087-91 are peptides derived from LSTRA and EL-4 tumours of Balb/c
CC mice. The peptides were tested for MHC specificity to find MHC I
CC specific peptides. These peptides are useful for tissue typing or for
CC screening for molecules that interact with MHC class I molecules. MHC
CC class I molecules can be identified using the peptides and also the
CC peptides are useful in vaccines against disease and infection e.g.
CC caused by viruses, bacteria or tumours.
CC
XX
SQ Sequence 8 AA;

Query Match 100.0%; Score 38; DB 18; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEKL 8
DB 1 STINFEKL 8

RESULT 6
AAW04642
ID AAW04642 standard; peptide; 8 AA.
XX
XX AAW04642;
AC
XX
XX 01-AUG-1997 (first entry)
DT
XX
XX Ovalbumin-derived activated CD8+ T cells epitope OVA8.
DE
XX
XX Macrophage; artificial antigen presenting cell; APC; cancer;
KM tumours; neoplasia; viral infection; retroviral infection;
KM autoimmune.
XX
XX Synthetic.
OS
XX
XX WO9637107-A1.
PN
XX
XX 28-NOV-1996.
PD
XX
XX 22-MAY-1996; 96WO-US07436.
PF
XX
XX 23-MAY-1995; 95US-0447761.
PR
XX
XX

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PA (SCRI ) SCRIPPS RES INST.
XX
XX DeBrujn MLH, Jackson MR, Peterson PA;
PI
XX WPI; 1997-020850/02.
DR
XX
XX Prodn. of activated CD8+ T cells directed to specific antigen - can
PT specifically kill target cells useful to treat, e.g. cancer
PT
XX
XX Example 1; Page 26; 84pp; English.
PS
XX
XX The method for the production of activated CD8+ T cells specifically
CC directed towards a particular antigen involves affixing peptides
CC corresponding to the particular antigen to an artificial support;
CC contacting macrophages with the affixed peptides for a time sufficient
CC for the peptides to be engulfed, and at least a portion of the peptides
CC to be presented on the surface of the macrophage; and contacting
CC unprimed CD8+ T cells with the peptide presenting macrophages for a
CC time sufficient to activate the unprimed CD8+ T cells. The present
CC sequence represents a peptide designated OVA8 which corresponds to
CC ovalbumin, a Kb-restricted peptide antigen. This is the optimal
CC peptide. Small extensions to the optimal peptide affect the affinity
CC of the peptide for soluble class I molecules in vitro e.g. the addition
CC of two amino acids to the amino-terminus lowers the affinity to Kb by
CC 76-fold compared to the optimal peptide; addition of two amino acids to
CC the carboxy-terminus lowers the affinity by 4-fold. The method,
CC macrophages and artificial antigen presenting cell, having a peptide
CC corresponding to the particular antigen present on its surface and at
CC least a portion of an artificial support in its interior, can be used to
CC treat conditions (e.g. cancer, tumours, neoplasia, viral or retroviral
CC infection or autoimmune or autoimmune-type conditions) in patients via
CC the specific killing of target cells.
XX
SQ Sequence 8 AA;

Query Match 100.0%; Score 38; DB 18; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEKL 8
DB 1 STINFEKL 8

RESULT 7
AAW80296
ID AAW80296 standard; Peptide; 8 AA.
XX
XX AAW80296;
AC
XX
XX 08-JAN-1999 (first entry)
DT
XX
XX Amino acids 257-264 of chicken ovalbumin used as an antigen.
DE
XX
XX Antisense oligonucleotide; antigen processing protein; TAP;
KM transporter; proteasome; antigen-presenting cell; cancer; infection;
KM cytotoxic T cell; chicken ovalbumin.
XX
XX Synthetic.
OS
XX
XX Gallus sp.
PN
XX
XX US5831068-A.
PD
XX
XX 03-NOV-1998.
PF
XX
XX 20-AUG-1996; 96US-0700035.
PR
XX
XX 20-AUG-1996; 96US-0700035.
PR
XX
XX 21-AUG-1995; 95US-0517373.
XX
XX (UYDU-) UNIV DUKE.
PA
XX
XX Gilboa E, Nair SK;
PI

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XX DR WPI; 1998-609331/51.
 XX PT Increasing the presentation of a peptide on a mammalian cell for
 XX production of antigen-presenting cells and stimulation of immune
 PT response - by contacting cells with antigen after inactivating the
 PT protein transporter associated with antigen processing or proteasome
 XX PS Disclosure; Column 12; 27pp; English.
 XX CC AAM80296-99 represent peptide antigens used in the course of the
 CC invention. The specification describes a method for increasing the
 CC presentation of a peptide (antigen) on a mammalian cell. The method
 CC comprises inhibiting the activity of a transporter associated with
 CC TAP or proteasome in the cell in vitro before contacting the cell
 CC with the peptide. Antigen-presenting cells produced as above can be used
 CC to stimulate an immune response in vitro or in vivo e.g. to treat or
 CC prevent cancer or infection with a pathogen, e.g. a bacterium or virus.
 CC Cytotoxic T cells produced as above can also be used for therapy.
 CC XX
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 19; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 1 SIINPEKL 8
 Db 1 SIINPEKL 8
 RESULT 8
 AAM70375
 ID AAM70375 standard; Protein; 8 AA.
 AC AAM70375;
 XX 18-NOV-1998 (first entry)
 DT
 XX Ovalbumin peptide used in the method of the invention.
 DE
 XX Ovalbumin; hep70; heat shock protein; vaccine; tumour therapy.
 KM
 XX Synthetic.
 OS
 OS Gallus sp.
 XX WO9835705-A1.
 PN 20-AUG-1998.
 XX 18-FEB-1998; 98WO-US03033.
 PF 25-NOV-1997; 97US-0066288.
 XX 18-FEB-1997; 97US-0038059.
 PR
 XX (MHED) WHITEHEAD INST BIOMEDICAL RES.
 PA
 XX Young RA;
 PI WPI; 1998-456872/39.
 XX Use of heat shock protein - for delivery of moiety into cells,
 PT useful for vaccination against tumours
 PT
 XX Disclosure; Fig 1A; 45pp; English.
 PS
 XX The present sequence represents residues 258-276 of the ovalbumin
 CC protein. The ovalbumin peptide was used in the method of the
 CC invention. The invention provides a method for delivering a moiety
 CC (e.g. ovalbumin protein) of interest into a cell which involves
 CC contacting the cell with a complex comprising the moiety of interest
 CC covalently linked to a heat shock protein (e.g. hep70). The method
 CC is claimed to be useful for providing the efficient delivery into cells

CC of moieties which are not normally able to enter cells or which enter
 CC cells only to a limited extent. The method can be useful for delivering
 CC moieties such as proteins, peptides, lipids, glycoproteins, small
 CC organic molecules and other molecules, particularly chemicals and other
 CC molecules which are useful therapeutically or diagnostically. The
 CC method is claimed to be useful when applied to vaccination regimes,
 CC e.g. for tumour therapy.
 CC XX
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 19; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Oy 1 SIINPEKL 8
 Db 1 SIINPEKL 8
 RESULT 9
 AAM68308
 ID AAM68308 standard; peptide; 8 AA.
 AC AAM68308;
 XX 25-MAR-2003 (updated)
 DT 14-OCT-1998 (first entry)
 XX MHC binding peptide Ova8.
 DE
 XX Antigen; major histocompatibility complex; MHC; lymphocyte; detection;
 KW immobilisation; cytotoxic T-cell; tumour; leukaemia; lymphoma;
 KW viral infection.
 XX Synthetic.
 OS
 OS WO9744667-A2.
 XX 27-NOV-1997.
 PD
 XX 21-MAY-1997; 97WO-FR00892.
 PF 21-MAY-1996; 96US-0651925.
 PR
 XX (INRM) INSERM INST NAT SANTE & RECH MEDICALE.
 PA (INSP) INST PASTEUR.
 XX Langlademoyen P, Lone Y, Kourilsky P, Abastado J;
 PI WPI; 1998-018653/02.
 XX Detection, purification and elimination of antigen-specific
 PT lymphocytes - for producing cytotoxic T cells for immuno-therapy of
 PT cancers and viral infection
 PT
 XX Disclosure; Page 24; 222pp; French.
 PS
 XX Peptides AAM68301-W68384 are examples of antigens (Ag) which can be
 CC loaded onto recombinantly produced major histocompatibility complex
 CC (MHC) molecules in a method of detecting antigen-specific lymphocytes.
 CC The MHC-antigen complex is then immobilised on a solid support and a
 CC sample containing cells recognising the MHC-Ag complex may be isolated.
 CC A similar method is used to isolate, purify or eliminate Ag-specific
 CC T-cells or to produce Ag-specific cytotoxic T-cells (CTC). The method
 CC is also used to detect and quantify tumour-specific T-cells and to
 CC generate CTC for specific killing of tumour cells (solid tumours,
 CC leukaemia or lymphoma) by injection into a human or animal, but also
 CC (Updated on 25-MAR-2003 to correct PI field.)
 CC XX
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 19; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEKL 8
| | | | |
DB 1 SIINFEKL 8

RESULT 10

AAW68365
ID AAW68365 standard; peptide; 8 AA.

AC AAW68365;

XX 25-MAR-2003 (updated)

DT 14-OCT-1998 (first entry)

DE MHC binding peptide from ovalbumin.

XX Antigen; major histocompatibility complex; MHC; lymphocyte; detection;
KW immobilisation; cytotoxic T-cell; tumour; leukaemia; lymphoma;
KM viral infection.

XX Synthetic.

PN MO9744667-A2.

PD 27-NOV-1997.

XX 21-MAY-1997; 97WO-FR00892.

XX 21-MAY-1996; 96US-0651925.

PA (INRM) INSERM INST NAT SANTE & RECH MEDICALE.

PA (INSP) INST PASTEUR.

PI Langlademeyen P, Lone Y, Kourilsky P, Abastado J;

DR WPI; 1998-018653/02.

PT Detection, purification and elimination of antigen-specific
PT lymphocytes - for producing cytotoxic T cells for immuno-therapy of
PT cancers and viral infection
XX
PS Disclosure; Page 29; 222pp; French.

CC Peptides AAW68301-W68384 are examples of antigens (Ag) which can be
CC loaded onto recombinantly produced major histocompatibility complex
CC (MHC) molecules in a method of detecting antigen-specific lymphocytes.
CC The MHC-antigen complex is then immobilised on a solid support and a
CC sample containing cells recognising the MHC-Ag complex may be isolated.
CC This peptide is derived from amino acids 258-276 of ovalbumin. A
CC similar method is used to isolate, purify or eliminate Ag-specific
CC T-cells or to produce Ag-specific cytotoxic T-cells (CTC). The method is
CC also used to detect and quantify tumour-specific T-cells. The method is
CC CTC for specific killing of tumour cells (solid tumours, leukaemia or
CC lymphoma) by injection into a human or animal, but also for treating
CC viral infections.
CC (Updated on 25-MAR-2003 to correct PI field.)
CC
XX
SQ Sequence 8 AA;

Query Match 100.0%; Score 38; DB 19; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEKL 8
| | | | |
DB 1 SIINFEKL 8

RESULT 11

AAW60700

ID AAW60700 standard; peptide; 8 AA.

XX AAW60700;

XX 22-SEP-1998 (first entry)

DE Ovalbumin peptide Tc1 peptide epitope (residues 257-264).

XX Immunisation; target antigen; epitope; inoculation; infant mammal;
KW viral antigen; depressed humoral response; respiratory syncytial virus;
KW rotavirus; measles virus; human immunodeficiency virus; hepatitis virus;
KW herpes simplex virus; influenza virus; Streptococcus pneumoniae;
KW Hemophilus influenzae; Neisseria meningitidis; Staphylococcus aureus;
KW protozoan antigen; malaria.

XX Unidentified.

OS WO9822145-A1.

PN 28-MAY-1998.

PD 21-NOV-1997; 97WO-US21687.

XX 22-NOV-1996; 96US-0755034.

PA (MOUN) MOUNT SINAI SCHOOL MEDICINE.

PI Bona C, Bot A;

DR WPI; 1998-312182/27.

PT Immunisation of infant mammals - by inoculating the mammal with a
PT nucleic acid encoding a relevant epitope of a target antigen
XX
PS Disclosure; Page 10; 83pp; English.

XX Sequence shown in AAW60683 to AAW60700 are epitope sequences of various
XX viral antigens used to exemplify the method of invention of immunising
XX an infant mammal against a target antigen. The method comprises
XX inoculating the mammal with a nucleic acid encoding a relevant epitope
XX of a target antigen in a carrier, such that the relevant epitope is
XX expressed in the infant mammal. The genetic immunisation of infant
XX mammals can give rise to effective cellular (including the induction of
XX cytotoxic T lymphocytes) and humoral immune responses against the target
XX antigen. The methods are particularly used for treating infants with
XX depressed humoral responses, that have high-zone tolerances against the
XX target antigens or have a Th2 biased immune response. The target antigen
XX may be a viral antigen, e.g. a respiratory syncytial virus antigen, a
XX rotavirus antigen, a measles virus antigen, a human immunodeficiency
XX virus antigen, a hepatitis virus antigen, a hepatitis B virus antigen, a
XX herpes simplex virus antigen or an influenza virus antigen, a bacterial
XX antigen e.g. Streptococcus pneumoniae antigen, Hemophilus influenzae
XX antigen, Neisseria meningitidis antigen, Staphylococcus aureus antigen
XX or a protozoan antigen such as a malaria antigen.

SQ Sequence 8 AA;

Query Match 100.0%; Score 38; DB 19; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEKL 8
| | | | |
DB 1 SIINFEKL 8

RESULT 12

AAW54265

ID AAW54265 standard; peptide; 8 AA.

XX AAW54265;

DT 30-JUL-1998 (first entry)

XX Ovalbumin peptide OVA.
 DE Pep-MHC complex; cytotoxic; T cell; cancer; ovarian; brain;
 KW major histocompatibility complex.
 XX Synthetic.
 OS
 XX MO9807441-A1.
 PN
 XX 26-FEB-1998.
 PD
 XX 22-AUG-1997; 97WO-US14814.
 PF
 XX 23-AUG-1996; 96US-0023437.
 PR
 XX (MASI) MASSACHUSETTS INST TECHNOLOGY.
 PA (UNITI) UNIV ILLINOIS FOUNO.
 XX
 PI Eisen HN, Kranz DM;
 XX WPI; 1998-168897/15.
 DR
 XX Composition for targeting an allo-reactive response to specific
 PT cells - comprises MHC-peptide complex bound to specific targeting
 PT agent, the complex having at least one allogenic component,
 PT specifically for treating ovarian and brain cancers
 XX
 PS Examples; Page 26; 45pp; English.
 XX
 CC Peptide OVA derived from ovalbumin was used in the production of soluble
 CC pep-MHC complexes. The pep-MHC (peptide-major histocompatibility
 CC complex) complexes, including both MHC heavy and light chains, bound to a
 CC specific targeting molecule can be used to target a cytotoxic T cell
 CC response to specific cells. This is particularly useful for targeting
 CC cancer cells, specifically of the ovary or brain. The method can also
 CC be used to eliminate an entire cell type for example during bone marrow
 CC therapy.
 CC
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 19; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEKL 8
 DB 1 SIINFEKL 8
 RESULT 13
 AAW52838
 ID AAW52838 standard; peptide; 8 AA.
 AC AAW52838;
 XX
 XX 24-JUN-1998 (first entry)
 DT
 XX Ovalbumin artificial target antigen.
 DE
 XX Ovalbumin; antigen; ATA; cytotoxic T lymphocyte; CTL; tumour; prevention.
 KW
 XX Gallus sp.
 OS
 XX MO9800163-A1.
 PN
 XX 08-JAN-1998.
 PD
 XX 18-JUN-1997; 97WO-US10195.
 PF
 XX 28-JUN-1996; 96US-0675332.
 PR
 XX (DAND) DANA FARBER CANCER INST INC.
 PA

PA (UYPI-) UNIV PITTSBURGH.
 XX
 XX Falo LD, Rock KL;
 PI
 XX WPI; 1998-086733/08.
 DR
 XX Inducing anti-tumour cytotoxic T lymphocytes - by cross-priming
 PT with artificial antigen, then immunisation with tumour cells
 PT modified in vitro to express the same antigen, does not require
 PT characterisation of tumour-specific antigens
 XX
 PS Example; Page 21; 44pp; English.
 XX
 CC Ovalbumin can be used as an artificial target antigen (ATA) to
 CC promote a cytotoxic T lymphocyte mediated response in mammals. Tumour
 CC cells from the host can be engineered to include ATA and therefore
 CC induce anti-tumour cytotoxic T lymphocytes. This method can be used for
 CC the treatment and prevention of a wide range of tumours even when
 CC the tumour is inaccessible or where metastases are being targeted.
 CC
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 19; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEKL 8
 DB 1 SIINFEKL 8
 RESULT 14
 AAY42307
 ID AAY42307 standard; peptide; 8 AA.
 AC AAY42307;
 XX
 XX 06-DEC-1999 (first entry)
 DT
 XX Ovalbumin-derived peptide antigen.
 DE
 XX Immunity; human leukocyte antigen; HLA; MHC; antigen;
 KW major histocompatibility complex; presentation; solubility;
 KW dendritic cell.
 XX
 OS Synthetic.
 XX
 XX MO9947646-A1.
 PN
 XX 23-SEP-1999.
 PD
 XX 19-MAR-1999; 99WO-US06627.
 PF
 XX 20-MAR-1998; 98US-0078832.
 PR
 XX (LUDW-) LUDWIG INST CANCER RES.
 PA (BIOP-) BIOPOLIO SCRL.
 XX
 XX Rescigno M, Girolomoni G, Corinti S, Ricciardi-Castagnoli P;
 PI WPI; 1999-571834/48.
 DR
 XX Preparation of dendritic cells which present antigens, used for
 PT stimulating an immune response by immunocompetent cells -
 PT
 PS Example 6; Page 13; 43pp; English.
 XX
 CC This sequence represents an ovalbumin-derived peptide antigen which
 CC is presented on major histocompatibility complex (MHC) Class I molecules
 CC of dendritic cells via the use of a novel process to improve soluble
 CC protein antigen presentation. This process uses dendritic cells which
 CC have internalised bacterial cells recombinantly expressing ovalbumin. The
 CC dendritic cells which have internalised the bacteria can be used to

CC generate an immune response which is stronger than the response which
 CC would be generated when soluble antigen alone is used. The dendritic
 CC cells which result from the internalisation are characterised by MHC
 CC Class I and Class II molecules which have much longer half lives and
 CC greater stability than comparable dendritic cells which have not
 CC internalised such bacteria. The dendritic cells can be contacted with
 CC immunocompetent cells for stimulating an immune response. The dendritic
 CC cells can also be used to stimulate maturation of an immature dendritic
 CC cell.

XX
 SQ Sequence 8 AA;

Query Match 100.0%; Score 38; DB 20; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 |||||
 DB 1 SIINFEKL 8

RESULT 15

AAV16871
 ID AAV16871 standard; peptide; 8 AA.

AC AAV16871;

DT 20-JUL-1999 (first entry)

XX OVA peptide sequence.

XX Conjugate peptide; heat shock protein; hsp; phage display library; virus;
 KM surface protein; tethering peptide; chaperone process; cytokine; cancer;
 KM neoplastic disease; infectious disease; bacterium; immune system; fungus;
 KM acquired immune deficiency; autoimmune disease.

XX Synthetic.

XX MO9922761-A1.

PN 14-MAY-1999.

XX 22-OCT-1998; 98WO-US22335.

XX 31-OCT-1997; 97US-0961707.

XX (SLOAN KETTERING INST CANCER RES.

PI Hartl U, Hoe MH, Houghton A, Mayhew M, Moroi Y;
 PI Querfeldt O, Rothman JF;

XX WPI; 1999-313177/26.

XX Identifying peptides which bind heat shock proteins

XX Examples; Page 49; 155pp; English.

CC The invention relates to conjugate peptides engineered to noncovalently
 CC bind to heat shock proteins (hsp). A method of identifying a hsp binding
 CC peptide comprises (a) contacting a phage display library having
 CC bacteriophage expressing, in a surface protein, inserted peptides with a
 CC hsp target, and bound to a benzquinone ansamycin antibiotic (BMA), in a
 CC physiologic binding buffer; (b) isolating a phage binding to the hsp
 CC target; and (c) identifying the inserted peptide expressed. The peptides
 CC which bind to a hsp can be used as tethering peptides for a hsp which may
 CC serve as an accessory in a chaperone process and/or may comprise a
 CC cytokine. They can also be coupled to antigens to induce an immune
 CC response. Such compositions can be used for treating neoplastic disease,
 CC e.g. cancers, infectious diseases, e.g. diseases caused by a bacterium,
 CC virus, protozoan, mycoplasma, fungus, yeast, parasite or prion, or a
 CC disease of the immune system, e.g. acquired immune deficiencies or
 CC autoimmune diseases.

SQ Sequence 8 AA;

Query Match 100.0%; Score 38; DB 20; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 |||||
 DB 1 SIINFEKL 8

RESULT 16

AAV03780
 ID AAV03780 standard; peptide; 8 AA.

AC AAV03780;

DT 23-JUN-1999 (first entry)

XX Ovalbumin peptide fragment (residues 257-264).

XX Dendritic cell-derived factor; proliferation; interferon gamma; IFNgamma;
 KM T cell; granulocyte macrophage colony-stimulating factor; GM-CSF;
 KM T cell stimulatory factor; lymphocyte; interleukin-2; lipopolysaccharide;
 KM autoimmune response; inflammatory; ovalbumin.

XX Synthetic.

XX Gallus gallus.

XX MO9918909-A2.

XX 22-APR-1999.

XX 14-OCT-1998; 98WO-US21614.

XX 14-OCT-1997; 97US-0062405.

XX (LUDWIG INST CANCER RES.

PI Dunn A, Marino MW, Noguchi Y, Old LJ, Wada H;

XX WPI; 1999-277418/23.

XX Dendritic cell and T cell derived factors for regulation of T cell

XX proliferation and interferon gamma production

XX Examples; Page 37; 68pp; English.

CC The invention relates to a dendritic cell-derived factor that restores
 CC proliferation and interferon gamma (IFNgamma) production to T cells from
 CC granulocyte macrophage colony-stimulating factor (GM-CSF). The dendritic
 CC cell-derived factor which is a T cell stimulatory factor modulates the
 CC effect of GM-CSF on production of IFNgamma by lymphocytes and the
 CC response of GM-CSF -/- T cells to interleukin-2, and corrects the
 CC lipopolysaccharide-induced defect in IFNgamma production. The dendritic
 CC cell-derived factor is used in vivo to increase proliferation of T cells
 CC and/or IFNgamma production, e.g. during immunisation to increase the
 CC response to an antigen. Agents with T cell derived factor activity are
 CC used to decrease production of IFNgamma, e.g. to reduce autoimmune
 CC responses and inflammatory reactions. Agents that bind to the factors,
 CC e.g. antisense sequences, are used to treat excessive/inadequate T cell
 CC proliferation and IFNgamma production. The dendritic cell-derived factor
 CC and T cell factor (or their fragments) can also be used to raise
 CC antibodies and as components in immunoassay and diagnostic systems, also
 CC for in vitro studies. The present sequence represents an ovalbumin
 CC peptide fragment used in the course of the invention.

XX Sequence 8 AA;

Query Match 100.0%; Score 38; DB 20; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 |||||
 KW 1 SIINFEKL 8

RESULT 17

AAW99480 standard; peptide; 8 AA.

AAW99480;

08-JUN-1999 (first entry)

Ovalbumin-derived peptide OVA.

Matrix protein; conjugate; mutant; major histocompatibility complex; MHC; class I molecule; beta-2-microglobulin; stimulation; immunity; tumour.

Synthetic.

WO9911775-A1.

11-MAR-1999.

20-AUG-1998; 98WO-US17308.

29-AUG-1997; 97US-0920413.

(GENO) GEN HOSPITAL CORP.

(HARD) HARVARD COLLEGE.

Garboczi DN, Walker J;

WPI; 1999-205182/17.

Method for conjugating a mutant major histocompatibility complex class I molecule and a compound - useful for stimulating immunity in an individual, and eradicating undesired cells, especially tumours

Example III-4; Page 22; 37pp; English.

The invention relates to the preparation of a conjugate of a mutant major histocompatibility complex (MHC) class I molecule (containing a Tyr67Cys amino acid substitution in the beta2-microglobulin subunit) and a compound. This sequence corresponds to the ovalbumin-derived peptide OVA. The peptide is used as a control to prepare a hybrid peptide-MHC class I tetramer in which a mutant human beta2-microglobulin subunit binds a mouse MHC class I molecule. The conjugates are useful for stimulating immunity in an individual, and eradicating undesired cells (e.g. tumours).

Sequence 8 AA;

Query Match 100.0%; Score 38; DB 20; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8

|||||
 DB 1 SIINFEKL 8

RESULT 18

AAW67585 standard; peptide; 8 AA.

AAW67585;

02-MAR-1999 (first entry)

T-cell activation peptide #3.

Activated T helper cell; CD4+; cytotoxic T cell; CD8+; liposome; epitope;

KW peripheral blood lymphocyte; antigen-presenting cell; APC; virus; tumour; bacterium; parasite; cytokine; vaccine; cancer; malaria; HIV; hepatitis; tuberculosis.

Synthetic.

WO9850527-A1.

12-NOV-1998.

07-MAY-1998; 98WO-US09288.

08-MAY-1997; 97US-0045949.

(BIOM-) BIOMIRA INC.

Agarwal B, Krantz MJ, Longenecker BM, Reddish MA;

WPI; 1999-034715/03.

Method of activation of T cells - by exposure to antigen-presenting cells loaded with antigen in liposome, used for, e.g. treating cancer and microbial infections

Disclosure; Page 6; 75pp; English.

Peptides AAW67583-W67611 are used to produce activated T helper (CD4+) and cytotoxic (CD8+) T cells. The activated T cells are produced by treating peripheral blood lymphocytes with liposome-encapsulated peptide antigen to generate Ag-loaded antigen-presenting cells (APC), contacting native or anergic T-cells with these APC, and isolating the resulting activated T-cells. The cells are specific for a particular antigen, particularly one derived from a tumour, but also those from viruses, bacteria and other parasites. It can also be used to identify antigens and epitopes able to generate an Ag-specific T-cell response (by assessing proliferation and cytokine release). Also the Ag-loaded APC can be used as cellular vaccines for treating cancer (claimed) or other diseases (e.g. malaria, human immune deficiency virus infection, hepatitis, tuberculosis). The activated T-cells can be used to treat the same conditions by adoptive T-cell transfer therapy.

Sequence 8 AA;

Query Match 100.0%; Score 38; DB 20; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8

|||||
 DB 1 SIINFEKL 8

RESULT 19

AAAB13763 standard; peptide; 8 AA.

AAAB13763;

02-FEB-2001 (first entry)

Peptide fragment from ovalbumin OVA.

T-cell; immune response; antigen; epitope; B7 family molecule; leukocyte function-associated antigen-3; LFA-3; intercellular adhesion molecule-1; ICAM-1; vaccine; immunotherapy; colon polyp; Crohn's disease; ulcerative colitis; breast lesion; tumour; ovalbumin.

Unidentified.

WO200034494-A1.

15-JUN-2000.

```

XX 12-NOV-1999; 99WO-US26866.
XX
XX 09-DEC-1998; 98US-0111582.
XX
XX (USSH) US DEPT HEALTH & HUMAN SERVICES.
XX (THER-) THERION BIOLOGICS CORP.
XX
XX Schlom J, Hodge J, Panicali D;
XX
XX WPI, 2000-431307/37.
XX
XX Novel recombinant vector useful as immunogens and vaccines for
XX stimulating and enhancing immunological responses to target cells and
XX antigens expresses multiple co-stimulatory molecules such as B7-1,
XX LFA-3, ICAM-1
XX
XX Example 31; Page 80; 188pp; English.
XX
XX Costimulatory molecules have important roles in T-cell activation and
XX therefore the immune response. The present invention relates to
XX recombinant vectors which comprise of foreign nucleic acid sequences
XX encoding at least three costimulatory molecules: a B7 family molecule,
XX leukocyte function-associated antigen-3 (LFA-3, human CD58) and
XX intercellular adhesion molecule-1 (ICAM-1, CD54) and optionally a foreign
XX gene encoding a target antigen or immunological epitope. The present
XX sequence is one such target antigen used in the present invention. The
XX present sequence is a tumour-associated antigen. The vector of the
XX present invention would be useful for providing an enhanced immune
XX response to the present target antigen. The vector of the present
XX invention may therefore be useful in immunotherapy for treating or
XX preventing diseases caused by viruses, bacteria, protozoans, parasites,
XX premalignant cells and tumour cells. The recombinant vector can be used
XX to treat or prevent preneoplastic or hyperplastic states such as colon
XX polyps, Crohn's disease, ulcerative colitis and breast lesions.
XX
XX Sequence 8 AA;
XX
XX Query Match 100.0%; Score 38; DB 21; Length 8;
XX Best Local Similarity 100.0%; Pred. No. 9.3e+05;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 STINPEKL 8
XX 1 STINPEKL 8
XX
XX DB 1 STINPEKL 8
XX
XX RESULT 20
XX AAB29465
XX ID AAB29465 standard; peptide; 8 AA.
XX
XX AC AAB29465;
XX
XX DT 09-FEB-2001 (first entry)
XX
XX DE Negative control peptide used in a cytotoxic T cell assay.
XX
XX KW Telomerase antigen variant; HLA-A2-binding; class I MHF;
XX human leukocyte antigen; major histocompatibility complex;
XX cytotoxic T-cell response; antigen-presenting cell; APC;
XX telomerase-expressing cell; cancer; anticancer vaccine.
XX
XX OS Synthetic.
XX
XX PN WO200061766-A2.
XX
XX PD 19-OCT-2000.
XX
XX PF 07-APR-2000; 2000WO-IB00610.
XX
XX PR 09-APR-1999; 99US-0128539.
XX
XX PA (BIOM-) BIOMIRA INC.

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XX Agrawal B, Longenecker BM;
XX
XX WPI, 2000-679493/66.
XX
XX New telomerase-specific T-cell antigens useful for generating T-cell
XX responses against telomerases and for producing vaccines for treating
XX or preventing cancer by in vivo or ex vivo techniques
XX
XX Example 1; Page 23; 34pp; English.
XX
XX The invention relates to a human telomerase peptide antigen (AAB29461)
XX which binds to a class I HLA (human leukocyte antigen, MHC, major
XX histocompatibility complex), and to conservatively substituted variants
XX thereof. The invention also relates to a vaccine comprising a telomerase
XX antigen or antigen variant, a nucleotide encoding a telomerase antigen
XX or variant, and a method of producing telomerase-primed antigen-
XX presenting cell (APC) comprising contacting an APC with a composition
XX containing a telomerase antigen or variant. The telomerase antigens or
XX vaccine compositions are useful for inducing a cytotoxic T-cell immune
XX response against telomerase and hence against telomerase-expressing
XX cells (i.e., cancer cells. Additionally, the telomerase antigen-primed
XX APC may be administered with interleukin-2 for cancer treatment or
XX prevention. The present sequence represents a peptide used in the
XX exemplification of the invention in an assay of the cytotoxic activity of
XX T-cells.
XX
XX Sequence 8 AA;
XX
XX Query Match 100.0%; Score 38; DB 21; Length 8;
XX Best Local Similarity 100.0%; Pred. No. 9.3e+05;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 STINPEKL 8
XX 1 STINPEKL 8
XX
XX DB 1 STINPEKL 8
XX
XX RESULT 21
XX AAB26484
XX ID AAB26484 standard; peptide; 8 AA.
XX
XX AC AAB26484;
XX
XX DT 16-JAN-2001 (first entry)
XX
XX DE Tumour associated OVA peptide.
XX
XX KW Immune response; vaccine; cancer; infection; tumour; OVA.
XX
XX OS Unidentified.
XX
XX PN WO200050080-A1.
XX
XX PD 31-AUG-2000.
XX
XX PF 23-FEB-2000; 2000WO-US04565.
XX
XX PR 26-FEB-1999; 99US-0261473.
XX
XX PA (UYDU-) UNIV DUKE.
XX
XX PI Gilboa E, Nair SK, Nicchitta CV;
XX
XX DR WPI, 2000-558368/51.
XX
XX PT Eliciting immune response in vertebrate for prevention and treatment of
XX cancer and infectious diseases involves administering purified complex
XX comprising calreticulin bound to an antigenic molecule
XX
XX PS Disclosure, Page 7; 82pp; English.
XX
XX The present invention relates to a method of eliciting an immune

```


CC response by administering a composition of a purified complex of
CC calreticulin bound to an antigenic molecule. The present invention is
CC useful for prevention and treatment of cancer and infectious disease
CC in a vertebrate especially of humans. The present sequence is the
CC tumour antigenic peptide OVA which was used in the invention.

XX
SQ Sequence 8 AA;

Query Match 100.0%; Score 38; DB 21; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEKL 8
DB 1 SIINFEKL 8

RESULT 22

AAB13785
ID AAB13785 standard; peptide; 8 AA.

XX AAB13785;

DT 10-NOV-2000 (first entry)

DE OVA-derived peptide.

KM Chicken; cytotoxic; vaccine; cytotoxic T cell; CTL; immunotherapy;
KM major histocompatibility complex class I; MHC class I; antigen; tumour;
KM prostate; breast; multiple myeloma; OVA peptide.

OS Gallus domesticus.

PN WO200035949-A1.

PD 22-JUN-2000.

PF 14-DEC-1999; 99WO-US29724.

PR 14-DEC-1998; 98US-0112324.

PA (DEND-) DENDREON CORP.

PI Laus R, Hakim I, Vidovic D;

DR WPI; 2000-442365/38.

PT Antigens modified by the covalent addition of a peptide that
PT facilitates entry into antigen presenting cells, useful for producing
PT compositions for immunizing against tumors and pathogens -

PS Example 1; Page 10; 34pp; English.

CC The present invention relates to compositions of modified soluble protein
CC antigens capable of eliciting an enhanced in vivo cytotoxic T cell (CTL)
CC response i.e. a major histocompatibility complex (MHC) class I molecule
CC response. The protein antigen is modified by the covalent addition of a
CC peptide sequence which facilitate entry of the antigen into antigen
CC presenting cells (APCs). The present sequence is a peptide derived from
CC the chicken antigen OVA. This peptide was used to prepare the modified
CC antigen. The modified antigen composition may be used for immunising
CC against, or treating a tumour e.g. prostate and breast carcinoma or
CC multiple myeloma, or pathogen in mammals.

XX
SQ Sequence 8 AA;

Query Match 100.0%; Score 38; DB 21; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEKL 8
DB 1 SIINFEKL 8

RESULT 23

AAV68311

XX AAV68311;

DT 13-APR-2000 (first entry)

DE Altered MHC determinant binding peptide SEQ ID NO:143.

KM MHC class I; major histocompatibility complex; microglobulin; antigen;
KM immune response; immunisation; AIDS; multiple sclerosis; toxic shock;
KM cancer; lupus erythematosus; snake bite; cytotoxic; antiviral;
KM immunomodulatory; dermatological; immunosuppressive; antiinflammatory;
KM neuroprotective.

OS Unidentified.

PN US6011146-A.

PD 04-JAN-2000.

PF 07-JUN-1995; 95US-0481985.

PR 15-NOV-1991; 91US-0792473.

PR 05-DEC-1991; 91US-0801818.

PA (INSP) INST PASTEUR.

PA (INRM) INST NAT SANTE & RECH MEDICALE.

PI Kourilsky P, Morteiz E, Abastado J;

DR WPI; 2000-125951/11.

PT New recombinant DNA encoding covalently linked form of major
PT histocompatibility complex Class I determinant, used for immune system
PT stimulation, e.g. for treating cancer -

PS Disclosure; Column 12; 88pp; English.

CC The present invention describes a recombinant DNA molecule (I)
CC containing a sequence (Ia) that encodes an altered MHC (major
CC histocompatibility complex) Class I determinant (II) comprises a
CC polypeptide with alpha1, alpha2, alpha3 and beta2-microglobulin
CC domains, in which alpha3 and beta2 are covalently linked, thorough C-
CC and N-terminal respectively, via a nucleotide spacer sequence encoding a
CC polypeptide. (II) includes an antigen-binding site and when (II) and
CC the antigen are associated they are recognized by a mammalian T cell
CC receptor (TCR). (I) are used to produce (II) which are used to study
CC functional interactions between the various MHC domains. They can also
CC be used to modulate (in vivo or in vitro) the immune system by inducing
CC an effector response (cytotoxicity, antibody synthesis, phagocytosis)
CC of immune system cells, typically for treating, or immunising against;
CC cancer, acquired immune deficiency syndrome, lupus erythematosus,
CC multiple sclerosis, toxic shock and snake bite, but also for selective
CC destruction of autoreactive cells, diagnostically to assay T cell
CC receptors and to raise specific antibodies (useful for diagnosis,
CC therapy, studying MHC-associated cellular processes and for affinity
CC purification). AA25758 and AAV68186 to AAV68316 are sequences used in
CC the exemplification of the present invention.

XX
SQ Sequence 8 AA;

Query Match 100.0%; Score 38; DB 21; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEKL 8
DB 1 SIINFEKL 8

```

RESULT 24
ID AAY59401
XX AAY59401 standard; Protein; 8 AA.
AC AAY59401;
XX
XX
XX 17-MAR-2000 (first entry)
XX
XX Ovalbumin protein fragment.
XX
XX Ovalbumin; chicken; nonvirulent bacterium; cytolysin;
XX diagnosis; gene therapy; polypeptide delivery; gene delivery.
XX
XX Gallus sp.
XX
XX US6004815-A.
XX
XX 21-DEC-1999.
XX
XX 13-AUG-1998; 98US-0133914.
XX
XX 13-AUG-1998; 98US-0133914.
XX
XX (REGC ) UNIV CALIFORNIA.
XX
XX Portnoy DA, Higgins DE;
XX
XX WPI; 2000-072064/06.
XX
XX Nonvirulent bacterium useful for the intracellular delivery of agents
XX to eukaryotic cells -
XX
XX Example; Column 12; 14pp; English.
XX
XX This sequence represents a fragment of the chicken ovalbumin.
XX The invention relates to a nonvirulent bacterium (1), comprising
XX a first gene encoding a nonsecreted foreign functional cytolysin operably
XX linked to a heterologous promoter which expresses cytolysin in the
XX bacterium and a second gene encoding a different foreign agent. The
XX bacterium can be used to deliver a wide variety of foreign agents to
XX eukaryotic cells for applications such as diagnosis, therapy including
XX prophylactic treatments such as immunisations and gene therapy especially
XX for single gene disorders suitable for localised treatment and
XX bioenhancement. They can be used to deliver antigenic polypeptides
XX presented in association with major histocompatibility (MHC) proteins to
XX antigen-presenting cells. The bacterium can be engineered to deliver
XX libraries of agents for screening. The foreign agent can be delivered to
XX any target cell capable of carrying out endocytosis of the subject
XX microbe, in particular epithelial cells, endothelial cells, muscle cells,
XX liver cells, pancreatic cells, neural cells, fibroblasts, tumour cells
XX and leukocytes. The foreign agent introduced into the cell is a nucleic
XX acid and/or protein which is bioactive in and therapeutic to the target
XX eukaryote. Bacterium mediated delivery of protein does not require
XX protein purification and high levels of protein can be delivered to the
XX cytosol of virtually all cells in culture. The process is rapid and
XX efficient and large enzymatically active proteins can be delivered.
XX
XX Sequence 8 AA;
XX
XX Query Match 100.0%; Score 38; DB 21; Length 8;
XX Best Local Similarity 100.0%; Pred. No. 9.3e+05;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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AC AAY52564;
XX
XX 28-FEB-2000 (first entry)
XX
XX Murine ovalbumin MHC class I epitope.
XX
XX Chimeric; ovalbumin; pan DR epitope; expression vector;
XX promoter; major histocompatibility complex; MHC; targeting; peptide;
XX epitope; antigen; presentation; class I; cytosolic pathway;
XX endoplasmic reticulum; class II; extracellular antigen;
XX endocytic pathway; helper T lymphocyte; HTL; universal epitope;
XX cytotoxic T lymphocyte; CTL; immune response; immunogenicity; assay;
XX vaccine; immunity; infection; pathogen; virus; HIV; HBV; HCV;
XX hepatitis B; hepatitis C; bacterium; protozoan; tumour cell;
XX autoimmune disease; activation; antiviral; antimetastatic;
XX immunoprotective.
XX
XX Synthetic.
XX
XX Mus sp.
XX
XX WO958658-A2.
XX
XX 18-NOV-1999.
XX
XX 13-MAY-1999; 99WO-US10646.
XX
XX 13-MAY-1998; 98US-0078904.
XX
XX 15-MAY-1998; 98US-0085751.
XX
XX (EPTM-) EPTMONE INC.
XX
XX Fikes JD, Hermanson GG, Sette A, Ishioka GY, Livingston B;
XX
XX Chestnut RW;
XX
XX WPI; 2000-039103/03.
XX
XX N-PSDB; AA238683.
XX
XX Expression vectors encoding major histocompatibility targeting
XX sequence, used as, e.g. tumor vaccines -
XX
XX Example 1; Page 45; 130pp; English.
XX
XX This sequence represents a murine ovalbumin MHC class I epitope
XX encoded by the AOS minigene insert of the expression vector PM1N.0
XX (AA238634). This insert encodes several MHC class I epitopes,
XX including this sequence, plus DNA encoding the universal MHC
XX class II (helper T) epitope, pan DR epitope (PADRE), and was used
XX in an exemplification of the present invention. The invention
XX relates to a novel expression vector comprising a promoter operably
XX linked to a fusion gene encoding a major histocompatibility complex
XX (MHC) targeting sequence, and two or more heterologous peptide
XX epitopes. The MHC targeting sequence may be a class I targeting
XX sequence, which directs an MHC class I epitope to a cytosolic pathway or
XX to the endoplasmic reticulum, or an MHC class II targeting sequence,
XX which directs extracellular antigens to enter the endocytic pathway to
XX be processed into antigen peptides for presentation on MHC class II
XX molecules. The heterologous epitopes may comprise either helper T
XX lymphocyte (HTL) epitopes, or a cytotoxic T lymphocyte (CTL) epitope and
XX a universal HTL epitope such as a pan DR epitope (PADRE). The vectors
XX are useful for stimulating an immune response in vivo, as well as for use
XX in assaying the human immunogenicity of a human T cell peptide epitope
XX in vivo in a non-human mammal. They provide a nucleic acid vaccine for
XX enhancing immunity against infectious pathogens, such as viruses (e.g.,
XX HIV, hepatitis B (HBV) and hepatitis C (HCV)) bacteria, protozoa (e.g.,
XX Plasmodium falciparum, the cause of malaria) and also tumour cells and
XX autoimmune diseases. Universal MHC class II epitopes are advantageously
XX combined with other MHC class I and class II epitopes to increase the
XX number of cells that are activated in response to a given antigen and
XX provide a broader population coverage of MHC-reactive alleles.
XX
XX Sequence 8 AA;
XX
XX Query Match 100.0%; Score 38; DB 21; Length 8;

```

Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINPEKL 8
1 SIINPEKL 8

Db 1 SIINPEKL 8

RESULT 26
AAV52965
ID AAV52965 standard; Peptide; 8 AA.

AC AAV52965;

DT 14-FEB-2000 (first entry)

DE Altered MHC determinant binding peptide SEQ ID NO:143.

XX Major histocompatibility complex; MHC class I; MHC class II; antigen;
KW immune response; diagnosis; antibody; immunisation; autoimmune disease;
KW acquired immune deficiency syndrome; AIDS; cytotoxic; dermatological;
KW anti-inflammatory; neuroprotective; immunosuppressive; antithyroid;
KW vaccine; lupus erythematosus; multiple sclerosis; thyroiditis;
KW toxic shock; tumour; snakebite.

OS Synthetic.

XX US5976551-A.

PD 02-NOV-1999.

PF 07-JUN-1995; 95US-0484905.

PR 05-DEC-1991; 91US-0801818.
PR 15-NOV-1991; 91US-0792473.

XX (INSP) INST PASTEUR.

PA (INRM) INSERM INST NAT SANTE & RECH MEDICALE.

PI Kourilsky P, Mottez E, Abastado J;

DR WPI; 2000-037081/03.

PT Composition containing an antigen and altered major histocompatibility
PT Class II determinant, used to immunize against autoimmune diseases,
PT e.g. acquired immune deficiency syndrome

PS Claim 8; Column 13; 96pp; English.

XX The present invention describes a composition capable of eliciting
CC anti-major histocompatibility (MHC) antibodies. The composition
CC comprises an antigen associated with an altered MHC Class II determinant
CC (I) comprising alpha1, alpha2, beta1 and beta2 polypeptide domains
CC encoded by a mammalian MHC Class II locus covalently linked to form a
CC polypeptide (I) containing beta2, alpha2, alpha1 and beta1 domains in
CC sequence. The resulting Antigen-MHC complex is recognizable by the T cell
CC receptor. The compositions are used for immunisation against, or
CC treatment of, a wide range of autoimmune diseases, e.g. acquired immune
CC deficiency syndrome (AIDS), lupus erythematosus, multiple sclerosis,
CC thyroiditis, toxic shock, tumour and snakebite, depending on the nature
CC of antigen. (I) is also used to analyse functional interactions between
CC the various domains and for targeting lymphocyte receptors. Antibodies
CC against (I) are produced by usual methods of immunisation or cell fusion,
CC and may be humanised by standard methods. These antibodies are useful for
CC diagnosis (detection or purification of MHC gene products), therapy
CC (neutralising MHC on cell surfaces) and in the study of MHC and cellular
CC processes. AA333240 to AA333242 and AAV52840 to AAV52970 represent
CC sequences used in the exemplification of the present invention.

XX Sequence 8 AA;

Query Match 100.0%; Score 38; DB 21; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINPEKL 8
1 SIINPEKL 8

Db 1 SIINPEKL 8

RESULT 27
AAG77871
ID AAG77871 standard; Protein; 8 AA.

AC AAG77871;

DT 08-MAY-2002 (first entry)

DE Ovalbumin 257-264 peptide.

XX Ovalbumin 257-264 peptide; major histocompatibility complex;
KW MHC-peptide complex; MHC; human; MHC class I alpha chain;
KW beta-2 microglobulin; MHC class II alpha chain; MHC class II beta chain;
KW vaccine; immune response modulation; hyperproliferative disorder;
KW neoplasm; hypergammaglobulinaemia; viral infection; hepatitis;
KW meningitis; bacterial infection; tuberculosis; gingivitis;
KW parasitic infection; autoimmune disease; Hashimoto's disease;
KW Graves' disease; rheumatoid arthritis; allergy; asthma; organ rejection;
KW graft-versus-host disease; GVHD; antigenic peptide.

OS Aves.

PN WO200178768-A2.

PD 25-OCT-2001.

PF 12-APR-2001; 2001WO-US11912.

PR 12-APR-2000; 2000US-196472P.

XX (UVRP) UNIV ROCHESTER.

PI Zauderer M, Smith ES;

DR WPI; 2001-602927/68.

PT Novel compound comprising major histocompatibility complex-peptide
PT complexes, used to modulate immune responses

PS Example 19; Page 101; 166pp; English.

XX The invention comprises a compound which contains one or more major
CC histocompatibility complex (MHC)-peptide complexes, and an antibody
CC specific for a cell surface marker. The complexes comprise an MHC class
CC I alpha chain, a beta-2 microglobulin molecule and an antigenic peptide
CC bound in the MHC groove. Alternatively, the complexes may comprise an MHC
CC class II alpha chain, an MHC class II beta chain, and an antigenic
CC peptide bound in the MHC groove. The complexes are linked to the carboxyl
CC termini of the antibody. The compounds of the invention can be used as a
CC vaccine to modulate an immune response. The compounds of the invention
CC are useful for treating hyperproliferative disorders (e.g. neoplasms and
CC hypergammaglobulinaemia); viral infections (e.g. hepatitis and
CC meningitis); bacterial infections (e.g. tuberculosis and gingivitis);
CC parasitic infections; autoimmune diseases (e.g. Hashimoto's disease,
CC Graves' disease and rheumatoid arthritis); allergic reactions/conditions
CC (e.g. asthma). The compounds of the invention may also be used in the
CC treatment of organ rejection or graft-versus-host disease (GVHD). The
CC present amino acid sequence represents the ovalbumin 257-264 peptide,
CC which was used as an antigenic peptide in an example of the invention.

XX Sequence 8 AA;

Query Match 100.0%; Score 38; DB 22; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Oy      1 SIINFEKL 8
XX      |||||
XX      1 SIINFEKL 8
Db

RESULT 28
AAM52562
ID AAM52562 standard; Peptide; 8 AA.
XX
XX AAM52562;
AC
XX 04-FEB-2002 (first entry)
XX
XX Cytotoxic T-cell epitope for ovalbumin.
DE
XX Cell death; toxic gene; tumour suppressor; ovalbumin;
XX Cytotoxic T-cell epitope.
XX
XX Unidentified.
OS
XX MO200172995-A2.
XX
XX 04-OCT-2001.
XX
XX 28-MAR-2001; 2001WO-US09953.
XX
XX 28-MAR-2001; 2000US-0192586.
XX 10-MAY-2000; 2000US-020343.
XX 23-JAN-2001; 2001US-0263226.
XX 27-FEB-2001; 2001US-0271426.
XX
XX (UNRP ) UNIV ROCHESTER.
XX
XX Zauderer M, Smith ES;
XX
XX WPI; 2001-570897/64.
XX
XX Selecting target polynucleotides, particularly toxic genes, involves
XX introducing a library of insert polynucleotides into a host cell
XX population, where the target polynucleotide promotes cell death -
XX
XX Example 1; Page 136; 359pp; English.
XX
XX The present invention relates to a method for selecting a target
XX polynucleotide. The method comprises introducing into a host cell
XX a population a library of insert polynucleotides, where expression of the
XX target polynucleotide directly or indirectly promotes host cell death.
XX The cells are cultured and the insert polynucleotides are collected from
XX the cells which die. The method is useful for selecting target
XX polynucleotides, particularly polynucleotides which alter cell phenotypes
XX of induce or inhibit cell death. The method can be used to isolate toxic
XX genes such as tumour suppressors. The present sequence was used to
XX illustrate the method of the the present invention.
XX
XX Sequence 8 AA;
SQ

Query Match      100.0%; Score 38; DB 22; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy      1 SIINFEKL 8
XX      |||||
XX      1 SIINFEKL 8
Db

RESULT 29
AAE13119
ID AAE13119 standard; peptide; 8 AA.
XX
XX AAE13119;
AC
XX 28-JAN-2002 (first entry)
XX

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XX DE Ovalbumin (OVA)-derived immunodominant octapeptide.
XX
XX Immunostimulatory fusion protein; IFP; antigen component; therapy;
XX immunostimulatory component; T-cell mediated immune response; DC;
XX dendritic cell; colon cancer; breast carcinoma; ovarian cancer;
XX Ovalbumin; OVA-derived immunodominant octapeptide.
XX
XX Unidentified.
OS
XX MO200174855-A2.
XX
XX 11-OCT-2001.
XX
XX 30-MAR-2001; 2001WO-US10515.
XX
XX 30-MAR-2000; 2000US-193504P.
XX
XX (DEND-) DENDREON CORP.
XX
XX Laus R, Vidovic D, Graddis T;
XX
XX WPI; 2001-662965/76.
XX
XX An immunostimulatory fusion protein comprising the intracellular domain
XX of HER-2 and an antigen elicits an immune response to the antigen and
XX is useful for the treatment of associated cancer associated -
XX
XX Example 1; Page 25; 59pp; English.
XX
XX The invention relates to immunostimulatory fusion proteins (IFP) and
XX nucleic acid molecules encoding such proteins. The IFPs comprise a
XX polypeptide antigen component and an immunostimulatory component derived
XX from the intracellular domain of HER-2 protein which is effective to
XX elicit a protective dendritic cell (DC)-induced T-cell mediated cellular
XX immune response to the antigen. IFP or superactivated dendritic cells
XX are used to treat cancer e.g. breast carcinoma, ovarian and colon cancer
XX associated with a particularly antigen. The present sequence is a
XX ovalbumin (OVA)-derived immunodominant octapeptide. This peptide
XX is used in the fusion constructs of the invention.
XX
XX Sequence 8 AA;
SQ

Query Match      100.0%; Score 38; DB 22; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy      1 SIINFEKL 8
XX      |||||
XX      1 SIINFEKL 8
Db

RESULT 30
AAE12145
ID AAE12145 standard; peptide; 8 AA.
XX
XX AAE12145;
AC
XX 15-JAN-2002 (first entry)
XX
XX Murine ovalbumin (OVA) peptide.
DE
XX Microbial delivery vehicle; prophylactic; immunisation; gene therapy;
XX tumour; carcinoma; neurodegeneration; muscular atrophy; cystostatic;
XX neuroprotective; antibacterial; insecticide; fungicide; antiviral;
XX antiprotocozal; cytostatic; anti-inflammatory; murine; ovalbumin; OVA;
XX listeriolysin O; LLO; MHC; major histocompatibility complex.
XX
XX Mus sp.
XX
XX US6287556-B1.
XX
XX 11-SEP-2001.
XX

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XX 21-DEC-1999; 99US-0469197.
XX
XX 13-AUG-1998; 98US-0133914.
XX
XX (REGC ) UNIV CALIFORNIA.
XX
XX Portnoy DA, Higgins DE;
XX
XX WPI; 2001-647179/74.
XX
XX Vaccine for preventing (e.g. as immunizations) or treating (e.g. as
XX gene therapy) tumor, carcinoma, neurodegeneration or muscular atrophy,
XX comprises a non-virulent bacterium
XX
XX Example; Column 12; 14pp; English.
XX
XX The invention relates to microbial-based intracellular delivery of
XX agents to eukaryotic cells. The agents include microbial delivery
XX vehicles such as nonvirulent bacteria comprising a first gene
XX encoding a nonsecreted foreign cyclolysin operably linked to a
XX heterologous promoter and a second gene encoding a different
XX foreign agent. The foreign agent may be a nucleic acid or protein,
XX and is frequently bioactive in and therapeutic to the target
XX eukaryote. The vaccine comprising nonvirulent bacterium is useful
XX for prophylactics (e.g. as immunisations) and treatments (e.g. as
XX gene therapy) of e.g. tumour, carcinoma, neurodegeneration or
XX muscular atrophy. The present sequence is murine ovalbumin (OVA)
XX peptide. This sequence is used to examine the ability of
XX Escherichia coli expressing listeriolysin O (LLO) and an antigenic
XX protein to deliver the antigen to the cytosol of macrophages for
XX processing and presentation on MHC (major histocompatibility
XX complex) class I molecules.
XX
XX Sequence 8 AA;
XX
XX Query Match 100.0%; Score 38; DB 22; Length 8;
XX Best Local Similarity 100.0%; Pred. No. 9.3e+05;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 SIINFEKL 8
XX |||||
XX 1 SIINFEKL 8
XX
XX DB
XX
XX RESULT 31
XX AAE09514
XX ID AAE09514 standard; peptide; 8 AA.
XX
XX AC AAE09514;
XX
XX DT 19-NOV-2001 (first entry)
XX
XX DE Human ovalbumin peptide.
XX
XX KM Mucin; cytosolic; immunostimulant; cell mediated immune response;
XX carcinoma; adenocarcinoma; breast cancer; dendritic cell; vaccine;
XX gene therapy; human; ovalbumin.
XX
XX OS Homo sapiens.
XX
XX PN WO200157068-A1.
XX
XX PD 09-AUG-2001.
XX
XX PF 01-FEB-2001; 2001WO-AU00090.
XX
XX PR 01-FEB-2000; 2000AU-0005369.
XX 14-JUN-2000; 2000US-0593870.
XX
XX (AUST-) AUSTIN RES INST.
XX
XX Mckenzie IFC, Pietersz GA, Apostolopoulos V;

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```

XX DR WPI; 2001-541537/60.
XX
XX Immunostimulant peptide, used as an anti-carcinoma vaccine, comprises a
XX an epitope of the non-VNTR, non-leader region of a mucin -
XX
XX Disclosure; Page 14; 84pp; English.
XX
XX The patent discloses peptide or polypeptides capable of eliciting
XX an immune response, comprising an amino acid sequence corresponding
XX to an epitope of the non-central portion of varying numbers of an
XX amino acid motif (VNTR), non-leader region of a mucin. The peptides
XX of the invention, fusion proteins comprising the peptide and conjugate
XX compounds with carbohydrate polymers are used to induce a cell mediated
XX immune response against mucin in the prevention or treatment of
XX carcinoma, preferably adenocarcinoma, most preferably breast cancer.
XX They are also used to pulse dendritic cell for in vivo transfer and
XX use as a vaccine. They are also used in gene therapy. The present
XX sequence is ovalbumin peptide from human. This sequence is used for
XX the prediction of T-cell epitopes.
XX
XX Sequence 8 AA;
XX
XX Query Match 100.0%; Score 38; DB 22; Length 8;
XX Best Local Similarity 100.0%; Pred. No. 9.3e+05;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 SIINFEKL 8
XX |||||
XX 1 SIINFEKL 8
XX
XX DB
XX
XX RESULT 32
XX AAG63855
XX ID AAG63855 standard; peptide; 8 AA.
XX
XX AC AAG63855;
XX
XX DT 29-OCT-2001 (first entry)
XX
XX DE Amino acid sequence of an OVA-derived minimal CTL peptide.
XX
XX KM Opi1; lipoprotein; adjuvant; type1 immune response; gp63;
XX leishmania major; leishmaniasis; TBC; leprosy; mycotin infection;
XX allergic asthma; autoimmune disease.
XX
XX OS Synthetic.
XX
XX PN WO200160404-A2.
XX
XX PD 23-AUG-2001.
XX
XX PF 13-FEB-2001; 2001WO-EP01673.
XX
XX PR 18-FEB-2000; 2000EP-0200589.
XX
XX (VLA-) VLAAMS INTERUNIVERSITAIR INST BIOTECHNOG.
XX
XX PI Revets H, Cornelis P, De Baetselier P;
XX
XX WPI; 2001-522552/57.
XX
XX Use of major Opi1 lipoprotein of Pseudomonas aeruginosa or its
XX functional fragments as adjuvant to obtain a Th1 type immune response
XX against heterologous antigen, for treating leishmaniasis, leprosy,
XX allergic asthma
XX
XX Disclosure; Page 12; 54pp; English.
XX
XX The present sequence represents an OVA-derived minimal CTL peptide,
XX which was used in the course of the invention. The specification
XX describes the use of the major Opi1 lipoprotein of Pseudomonas
XX aeruginosa or its functional fragments as an adjuvant to obtain

```

CC a Th1 type immune response against a heterologous antigen. They are
 CC especially used as an adjuvant to obtain a Th1 type immune response
 CC against a heterologous antigen such as antigen gp63 of *Leishmania* major,
 CC for treating a disease such as leishmaniasis, TB (undefined), leprosy,
 CC mycobacter infection, allergic asthma or an autoimmune disease, in which
 CC the natural Th1 response is insufficient and/or in which the immune
 CC response is polarizes towards Th2 response.

CC Sequence 8 AA;

Query Match 100.0%; Score 38; DB 22; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05; Mismatches 0; Indels 0; Gaps 0;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 STINPEKL 8
 |||||
 1 STINPEKL 8

RESULT 33

AA066422 standard; peptide; 8 AA.

AC AAG6422;

DT 23-OCT-2001 (first entry)

DE Chicken ovalbumin peptide, OVA257-264, used as a peptide antigen.

XX Immunomodulator; vaccine; immune response; immunogenic; chicken;

KW ovalbumin.

XX Gallus domesticus.

PN WO200154720-A1.

PD 02-AUG-2001.

PS 05-JAN-2001; 2001WO-EP00087.

PR 28-JAN-2000; 2000AT-0000129.

PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH.

PI Lingnan K, Mattnier F, Schmidt W, Birnstiel M, Buschle M;

DR WPI; 2001-536419/59.

XX Pharmaceutical composition useful for inducing immune response

PT comprises antigen, immunogenic oligodeoxynucleotide containing

PT cytosine-guanine dinucleotide motifs and polycationic polymer -

XX Example 1; Page 22; 39pp; English.

XX The present invention relates to a pharmaceutical composition which

CC comprises an antigen, an immunogenic oligodeoxynucleotide containing

CC cytosine-guanine dinucleotide (CpG) motifs (CpG-ODN) and a polycationic

CC polymer. The composition is useful for making a vaccine to induce potent

CC immune responses, or to decrease or ablate undesired immune responses.

CC The present sequence, OVA257-264, is a peptide from chicken ovalbumin.

CC This sequence was used as a peptide antigen in the method of the present

XX invention.

XX Sequence 8 AA;

Query Match 100.0%; Score 38; DB 22; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05; Mismatches 0; Indels 0; Gaps 0;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

1 STINPEKL 8
 |||||
 1 STINPEKL 8

RESULT 34

AA05398 standard; peptide; 8 AA.

AC AA05398;

DT 24-OCT-2001 (first entry)

DE Peptide released from ovalbumin (Ova) after cleavage of peptide PI.

XX Heat shock protein; hsp; CD8+ cytotoxic T lymphocyte; ovalbumin;

KW CTL; CD4+ T cell; AIDS; acquired immunodeficiency syndrome; murine;

KW human immunodeficiency virus; HIV; pathogen; cancer.

XX Mus sp.

PN WO200151081-A1.

PD 19-JUL-2001.

PS 01-DEC-2000; 2000WO-US32831.

PR 14-JAN-2000; 2000US-0176143.

PA (WHEB) WHITEHEAD INST BIOMEDICAL RES.

PI (MASI) MASSACHUSETTS INST TECHNOLOGY.

PI Huang Q, Richmond JFL, Cho BK, Palliser D, Chen J, Eisen HN;

DR WPI; 2001-451815/48.

PT Inducing a CD8+ cytotoxic T lymphocyte immune response in an individual

PT for treating diseases such as HIV involves administering a fusion

PT molecule comprising a heat shock protein -

XX Example 1; Fig 1C; 58pp; English.

XX The present sequence represents a naturally occurring murine peptide

CC which is released from ovalbumin (Ova) upon cleavage of peptide PI

CC (AA05397). The present sequence is described in an invention

CC relating to a novel method of inducing a CD8+ cytotoxic T lymphocyte

CC (CTL) response to a molecule in an individual by administering a

CC fusion molecule joined to a hsp, or an adenosine triphosphate (ATP)

CC binding domain of a hsp. The method is particularly useful in inducing

CC a CD8+ CTL response in an individual deficient in CD4+ T cells e.g. for

CC treating an AIDS acquired immunodeficiency syndrome patient carrying

CC the human immunodeficiency virus (HIV). The method is also useful for

CC treating diseases that are caused by or associated with intracellular

CC pathogens, and for treating cancer.

XX Sequence 8 AA;

Query Match 100.0%; Score 38; DB 22; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05; Mismatches 0; Indels 0; Gaps 0;

RESULT 35

AA06033 standard; peptide; 8 AA.

AC AA06033;

DT 25-SEP-2001 (first entry)

DE Chicken ovalbumin CTL epitope.

PT controlling immune responses -
 XX Example 6; Page 69; 97pp; English.
 CC The present invention describes immunomodulatory populations ((i) and
 CC ((ii)) of conjugate molecules (CMs) comprising immunostimulatory sequences
 CC ((ISS)) of polynucleotides and antigens. The extent of conjugation affects
 CC the immunological properties (e.g. the extent of antigen-specific
 CC antibody formation, including Th1-associated antibody formation) so the
 CC conjugates are used for altering the type and extent of immune response.
 CC ((i) and ((ii)) have immunomodulatory, immunosuppressive and anti-allergic
 CC activities, and can be used in the modulation of immune responses via
 CC the stimulation of Th1 lymphocytes and Th1-associated cytokines, and
 CC suppression of Th2 lymphocytes and cytokines. The populations ((i) and
 CC ((ii)) of conjugate molecules may be used for modulating immune responses
 CC in individuals e.g. for the treatment of an allergic condition. ((i) and
 CC ((ii)) may be used to modulate immune responses and therefore prevent
 CC potentially harmful reactions to antigens. The present sequence
 CC represents an ovalbumin (OVA) cytotoxic T lymphocyte (CTL) epitope
 CC which is used in the exemplification of the present invention.
 XX
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 22; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEKL 8
 DB 1 SIINFEKL 8
 RESULT 38
 ID AAB82176 standard; peptide; 8 AA.
 AC AAB82176;
 XX
 DT 20-JUL-2001 (first entry)
 DE Immunodominant CTL epitope of ovalbumin.
 XX
 KW Vaccine; Antiviral; Antibacterial; Antiparasitic; liposome;
 KW archaeobacterium; cytotoxic T lymphocyte response;
 KW immunodominant epitope; ovalbumin; archaeosome.
 XX
 OS Unidentified.
 XX
 PN WO200126683-A2.
 XX
 PD 19-APR-2001.
 XX
 PF 12-OCT-2000; 2000WO-CA01197.
 XX
 PR 12-OCT-1999; 99US-0158944.
 PR 08-JUN-2000; 2000US-0209988.
 XX
 PA (CANA) NAT RES COUNCIL CANADA.
 XX
 PI Sprott GD, Krishnan L, Conlan JW, Omri A, Patel GB;
 DR WPI; 2001-281839/29.
 XX
 PT New vaccine comprising a liposome useful for conferring protective
 PT immunity against an intracellular pathogen -
 XX
 PS Disclosure; Page 34; 98pp; English.
 CC The present invention relates to a vaccine composition comprising a
 CC liposome prepared from the total polar lipid extract of an
 CC archaeobacterium and an acellular antigen, preferably an isolated
 CC outer membrane from a pathogen. The vaccine of the invention provides an
 CC enhanced cytotoxic T lymphocyte response. The vaccine of the invention

CC is useful for conferring protective immunity against an intracellular
 CC pathogen. The present peptide: immunodominant CTL epitope of ovalbumin,
 CC was used to illustrate the present invention. This peptide was used to
 CC test for the ability of archaeosomes to induce CTL responses to
 CC ovalbumin.
 XX
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 22; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEKL 8
 DB 1 SIINFEKL 8
 RESULT 39
 ID AAB81122 standard; peptide; 8 AA.
 AC AAB81122;
 XX
 DT 04-JUL-2001 (first entry)
 DE Chicken ovalbumin (OVA) peptide.
 XX
 KW Ovalbumin; chicken; OVA; immune response; Langerhans cell migration;
 KW tumour; EGF-OVA.
 XX
 OS Gallus gallus.
 XX
 PN US6210672-B1.
 XX
 PD 03-APR-2001.
 XX
 PF 20-OCT-1998; 98US-0176044.
 XX
 PR 20-OCT-1998; 98US-0176044.
 XX
 PA (TORR-) TORREY PINES INST MOLECULAR STUDIES.
 XX
 PI Cowling C;
 DR WPI; 2001-280845/29.
 XX
 PT Enhancing an immune response to an antigen in a mammal comprises
 PT topically administering the antigen, a penetration enhancer and an
 PT agent for enhancing Langerhans cell migration -
 XX
 PS Disclosure; Column 4; 15pp; English.
 XX
 CC This invention relates to a method for enhancing an immune response to an
 CC antigen in a mammal. The method comprises administering a composition
 CC comprising the antigen, a penetration enhancer selected from lipophilic
 CC solvents, low-frequency ultrasound, electroporation, iontophoresis and
 CC intraepidermal delivery and an agent for enhancing Langerhans cell
 CC migration to an epidermal or mucosal site. The method can be used to
 CC enhance the immune response to tumours, viruses, bacteria and parasites.
 CC The present sequence represents a fragment of the chicken ovalbumin (OVA)
 CC protein. The peptide functions as a EGF-OVA tumour associated peptide
 CC antigen for CD8+ cytotoxic T lymphocytes. The peptide can be used in the
 CC method of the invention to enhance an immune response to the EGF-OVA
 CC tumour.
 XX
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 22; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEKL 8
 DB 1 SIINFEKL 8

DB 1 SIINEKL 8

RESULT 40
AAB82065 standard; peptide; 8 AA.

ID AAB82065 standard; peptide; 8 AA.
XX
XX AAB82065;
XX
XX 22-JUN-2001 (first entry)
XX
XX Ovalbumin-derived peptide, used as a control peptide.
XX
XX Antigen; immunostimulant; vaccine; pharmaceutical composition; antiviral;
XX
XX viral infection; ovalbumin.
XX
XX Unidentified.
XX
XX WO200124822-A2.
XX
XX 12-APR-2001.
XX
XX 02-OCT-2000; 2000WO-EP09657.
XX
XX 01-OCT-1999; 99AT-0001680.
XX
XX (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
XX
XX Fleitmann J, Mattner F, Buschle M, Melling J;
XX
XX WPI; 2001-290577/30.
XX
XX New pharmaceutical composition comprising an antigen, an
XX
XX immunostimulating substance and a polycationic polymer, useful in
XX
XX manufacturing vaccines -
XX
XX
XX Example 1; Page 11; 20pp; English.

CC The present invention relates to a pharmaceutical composition comprising
CC (a) an antigen; (b) an immunostimulating substance consisting of
CC neuroactive compounds, hormones, compounds having growth hormone activity
CC or their mixtures; and (c) a polycationic polymer. The composition is
CC useful in manufacturing vaccines. To illustrate the present invention, a
CC murine tyrosinase related protein-2 peptide (TRP-2 peptide; see
CC AAB82064), was used. Mice were injected subcutaneously with either the
CC TRP-2 peptide, TRP-2 peptide + human growth hormone (HGH), TRP-2 peptide
CC + poly-L-arginine 60 (pR60) or TRP-2 peptide + pR60 + HGH. Animals were
CC sacrificed 10 days post injection, and mesenteric and inguinal lymph
CC nodes were harvested. Lymphocytes were prepared from lymph nodes and were
CC re-stimulated with TRP-2 peptide or with an ovalbumin-derived peptide
CC (the present peptide), with the same major histocompatibility complex
CC (MHC) restriction serving as negative control. Spots representing single
CC T cells specific for the peptide used for re-stimulation were counted. No
CC spots were detected when the ovalbumin derived peptide was used, while
CC TRP-2 peptide + pR60 + HGH showed the highest number of spots or single T
CC cells. The present peptide was also used as a control peptide for
CC experiments with substance P (see AAB82070).
XX
XX
XX Sequence 8 AA;
SQ

Query Match 100.0%; Score 38; DB 22; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINEKL 8
DB 1 SIINEKL 8

RESULT 41
AAB92374 standard; Peptide; 8 AA.
XX

AC AAB92374;
XX
XX 22-JUN-2001 (first entry)
XX
XX Miscellaneous peptide SEQ ID NO:1550.
XX
XX
XX Protection; endogenous therapeutic peptide; peptidase; conjugation;
XX
XX blood component; modification; succinimidyl; maleimido group; amino;
XX
XX hydroxyl; thiol; hormone; growth factor; neurotransmitter.
XX
XX Homo sapiens.
XX
XX Synthetic.
XX
XX WO200069900-A2.
XX
XX 23-NOV-2000.
XX
XX 17-MAY-2000; 2000WO-US13576.
XX
XX
XX 17-MAY-1999; 99US-0134406.
XX
XX 10-SEP-1999; 99US-0153406.
XX
XX 15-OCT-1999; 99US-0159783.
XX
XX (CONJ-) CONJUCHEM INC.
XX
XX
XX Bridon DP, Ezrin AM, Milner PG, Holmes DL, Thibaudau K;
XX
XX WPI; 2001-112059/12.
XX
XX
XX Modifying and attaching therapeutic peptides to albumin prevents
XX
XX peptidase degradation, useful for increasing length of in vivo activity
XX
XX
XX Disclosure; Page 711; 733pp; English.

CC The present invention describes a modified therapeutic peptide (I)
CC comprising a therapeutically active amino acid region (Ii) and a
CC reactive group (Iii) (e.g. succinimidyl and maleimido groups) attached to
CC a less therapeutically active amino acid region (Iv), which covalently
CC bonds with amino/hydroxyl/thiol groups on blood components to form a
CC peptidase stabilised therapeutic peptide composed of 3-50 amino acids.
CC (Ii) are useful for modifying therapeutic peptides e.g. hormones, growth
CC factors and neurotransmitters, to protect them from peptidase activity
CC in vivo for the treatment of various disorders. Endogenous therapeutic
CC peptides are not suitable as drug candidates as they require frequent
CC administration due to rapid degradation by peptidases in the body.
CC Modifying and attaching therapeutic peptides to albumin prevents or
CC reduces the action of peptidases to increase length of activity (half
CC life) and specificity as bonding to large molecules decreases
CC intracellular uptake and interference with physiological processes.
CC AAB90829 to AAB92441 represent peptides which can be used in the
CC exemplification of the present invention.
XX
XX
XX Sequence 8 AA;
SQ

Query Match 100.0%; Score 38; DB 22; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINEKL 8
DB 1 SIINEKL 8

RESULT 42
AAB48950 standard; Protein; 8 AA.
XX
XX AAB48950;
XX
XX 27-MAR-2001 (first entry)
XX
XX Ovalbumin MHC class I epitope, SEQ ID NO:6.
XX

```

XX Transposable element; MHC epitope; major histocompatibility complex;
KW intracellular bacterial pathogen; loxp site; Cre recombinase;
KW insertion end; in-frame fusion; detection; antigen;
KW disseminated insertions of class-I epitopes; DICE-I; transposon Tn5;
KW ovalbumin MHC class I epitope.
XX Unidentified.
XX MO200071158-A1.
XX 30-NOV-2000.
XX 26-MAY-2000; 2000WO-US14687.
XX 26-MAY-1999; 99US-0136210.
XX (UYOR-) UNIV OREGON HEALTH SCI.
XX Heffron FL, Parker DC, Ellefsen DD;
XX WPI; 2001-031967/04.
XX Transposable element for detecting an antigenic epitope of a pathogen,
XX comprising 5' and 3' recombining sites, nucleic acid sequences encoding
XX a selectable marker and major histocompatibility complex (MHC) epitope,
XX and an insertion end -
XX Claim 5; Page 41; 63pp; English.
XX The invention relates to a novel transposable element comprising DNA
XX encoding a selectable marker (e.g., antibiotic resistance) located
XX between a 5' recombining site and a 3' recombining site (e.g., loxp
XX sites); DNA encoding an MHC (major histocompatibility complex) epitope
XX either 5' of the 5' recombining site or 3' of the 3' recombining site;
XX and insertion ends comprising an inverted repeat sequence at the 5' and
XX 3' ends of the transposable element sufficient for integration of the
XX transposable element. The transposable elements of the invention are able
XX to introduce in-frame insertions throughout the chromosome of an
XX intracellular bacterial pathogen. This system "tags" the bacterial gene
XX and resulting protein, allowing the identification of proteins
XX secreted across the membranes of the eukaryotic cell infected by the
XX bacterium. In one embodiment, the transposable elements contain an
XX antibiotic resistance cassette, two minimal loxp recombination sites, an
XX MHC class I or class II epitope, and flanking insertion ends. A
XX transposase, such as the Cre recombinase protein, is expressed in trans
XX from a plasmid, or can be included in the transposable element. The Cre
XX recombinase loops out the intervening sequences containing the antibiotic
XX resistance cassette. When the transposable element inserts within a gene,
XX the resolved insertion places the MHC class I or class II epitope in
XX frame with the gene. The transposable elements of the invention are
XX useful for detecting an antigenic epitope of an intracellular bacterial
XX pathogen, such as Salmonella sp., Mycobacterium tuberculosis and Listeria
XX monocytogenes. Certain embodiments of the technology, termed
XX "disseminated insertions of class-I epitopes" (DICE-I; DICE-II for
XX class II epitopes) allow the rapid and accurate identification of
XX proteins involved in bacterial pathogenesis so that such proteins can
XX be used as vaccine and drug targets. Carrier vaccines may be generated
XX by infecting bacteria with a transposable element of the invention
XX which additionally comprises an antigen associated with a disease,
XX preferably cancer or a viral or bacterial disease, operably linked to the
XX MHC epitope DNA of the transposable element. The present sequence
XX represents an ovalbumin MHC class I epitope specifically claimed
XX for use in the invention.
XX
SQ Sequence 8 AA;

```

```

Query Match          100.0%; Score 38; DB 22; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 SIINFEXL 8
|||||||
XX

```

```

DB 1 SIINFEXL 8
RESULT 43
AAE28959
ID AAE28959 standard; peptide; 8 AA.
XX
XX AAE28959;
XX
XX 27-JAN-2003 (first entry)
XX
XX Chicken ovalbumin peptide.
XX
XX Modified vaccinia Ankara virus; MVA; HIV; human immunodeficiency virus;
KW CD8+ T cell; immune response; acquired immune deficiency syndrome; AIDS;
KW viral infection; vaccine; immunostimulant; virucide; chicken.
XX
XX Gallus sp.
XX
XX WO200272754-A2.
XX
XX 19-SEP-2002.
XX
XX 01-MAR-2002; 2002WO-US06713.
XX
XX 08-MAR-2001; 2001US-274434P.
XX
XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
XX Moss B, Wyatt L, Earl P;
XX
XX WPI; 2002-723330/78..
XX
XX New recombinant modified vaccinia Ankara (MVA) virus expressing HIV
XX env, gag and pol genes, useful for boosting or inducing CD8 T cell
XX immune responses in primates, e.g. humans, particularly for preventing
XX AIDS or other viral infections -
XX
XX Example 1; Page 111; 112pp; English.
XX
XX The present invention relates to a composition comprising recombinant
XX modified vaccinia Ankara (MVA) virus expressing an HIV (human immuno-
XX deficiency virus) env, gag and pol gene or its modified gene for the
XX production of an HIV Env, Gag and Pol antigen by expression from the
XX recombinant MVA virus. The HIV env gene is modified to encode an HIV
XX Env protein composed of gp120 and the membrane-spanning and ectodomain
XX of gp41, but lacking part or all of the cytoplasmic domain of gp41. The
XX composition or recombinant MVA virus is useful for boosting or inducing
XX CD8+ T cell immune response in primates, particularly in humans. The
XX composition may be used for preventing AIDS (acquired immune deficiency
XX syndrome) or other viral infections. Sequences of the invention are also
XX used as vaccines. The present sequence is chicken ovalbumin peptide. This
XX sequence is used in the exemplification of the invention.
XX
SQ Sequence 8 AA;

```

```

Query Match          100.0%; Score 38; DB 23; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY 1 SIINFEXL 8
|||||||
DB 1 SIINFEXL 8

```

```

RESULT 44
AAE26368
ID AAE26368 standard; peptide; 8 AA.
XX
XX AAE26368;
XX
XX 13-DEC-2002 (first entry)
XX

```

```

DE      Ovalbumin CTL epitope.
XX
XX      Human, immune response, T-helper cell epitope; chitosan; CTL response;
KW      vaccine; prostate cancer; breast cancer; cytostatic; immunostimulant.
XX
XX      Unidentified.
XX
XX      WO200234287-A2.
XX
XX      02-MAY-2002.
XX
XX      26-OCT-2001; 2001WO-DK00705.
XX
XX      27-OCT-2000; 2000DK-0001606.
XX      03-NOV-2000; 2000US-245166P.
XX      18-JUN-2001; 2001DK-0000936.
XX
XX      (PHAR-) PHARMEXA AS.
XX
XX      Beier AM, Gautam A, Nouritsen S;
XX      WPI; 2002-463339/49.
XX
XX      Inducing or enhancing an immune response against an antigen,
XX      particularly cytotoxic T-lymphocyte responses, for treating or
XX      ameliorating prostate or breast cancer, comprises administering the
XX      antigen formulated with chitosan
XX
XX      Example 3; Page 63; 97pp; English.
XX
XX      The invention relates to a method for inducing or enhancing an immune
XX      response against a polypeptide antigen in an animal, including human.
XX      The method comprises administering the polypeptide antigen or at least
XX      one variant which includes at least one first T-helper cell epitope that
XX      is foreign to the animal (foreign TH epitope) and is formulated with
XX      chitosan. The polypeptide antigen is weakly immunogenic or non-
XX      immunogenic. The invention is used as vaccine. The chitosan and
XX      polypeptide antigen or its variant are useful in the preparation of an
XX      immunogenic composition for inducing or enhancing an immune response,
XX      particularly CTL response, against the polypeptide or protein antigen.
XX      The method for inducing or enhancing an immune response is useful in
XX      treating or ameliorating cancer, e.g. prostate or breast cancer. The
XX      present sequence is ovalbumin CTL epitope used to illustrate the method
XX      of the invention.
XX
XX      Sequence 8 AA;
XX
XX      Query Match          100.0%; Score 38; DB 23; Length 8;
XX      Best Local Similarity 100.0%; Pred. No. 9.3e+05;
XX      Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0.
XX
XX      QY      1 SIINFEKL 8
XX      DB      1 SIINFEKL 8
XX
XX      RESULT 45
XX      ABB79933
XX      ID      ABB79933 standard; Peptide; 8 AA.
XX
XX      AC      ABB79933;
XX
XX      DT      12-DEC-2002 (first entry)
XX
XX      DE      Ovalbumin T-cell epitope.
XX
XX      KW      Vaccine; genetic immunisation; gene therapy; antigen; epitope;
XX      T-cell; T-lymphocyte; ovalbumin.
XX
XX      OS      Unidentified.
XX
XX      PN      US2002115625-A1.
XX

```

PD 22-AUG-2002.
 XX 08-MAR-2001; 2001US-0801540.
 PF 19-MAY-1999; 99US-0308511.
 PR (BONA/) BOT A.
 PA (BONA/) BONA C.
 XX Bot A, Bona C;
 PI WPI; 2002-712482/77.
 DR
 XX Immunizing an infant mammal against a target antigen or inducing a
 PT cytotoxic T cell response against a pathogen in the mammal, comprises
 PT administering nucleic acid encoding relevant epitopes of pathogen
 PT associated target antigens -
 XX
 XX Disclosure; Page 4; 45pp; English.
 PS
 XX The present sequence is that of an ovalbumin T-cell epitope (amino
 CC acids 257-264). This is a T-CTL epitope which, in the context of
 CC MHC Class II self antigens, may be recognised by a cytotoxic T-cell
 CC and thereby promote CTL-mediated lysis of cells comprising the target
 CC antigen. It is an example of T-cell epitopes which may be used
 CC according to the invention. The invention relates to immunising an
 CC infant mammal against a target antigen or inducing a cytotoxic
 CC T-cell response against a pathogen. The method involves inoculating
 CC the infant with a nucleic acid encoding one or more relevant epitopes
 CC of one or more target antigens associated with the pathogen in a
 CC carrier, so that the relevant epitope(s) is expressed in the infant
 CC mammal. B- or T-cell epitopes may be used, and the pathogen may be
 CC a virus, bacterium, protozoan, fungus, yeast, or parasite. The method
 CC may reduce the need for subsequent boost administrations and may
 CC prevent the side-effects associated with live attenuated vaccines.
 CC Administration of multiple epitopes directed to antigens
 CC associated with more than one pathogen may provide an infant with a
 CC broader spectrum of protection, and may be a means for inducing an
 CC immune response to a variety of childhood pathogens.
 XX
 SQ Sequence 8 AA;
 QY 1 SIINFEKL 8
 DB 1 SIINFEKL 8
 Query Match 100.0%; Score 38; DB 23; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 RESULT 46
 ABG93028
 ID ABG93028 standard; Peptide; 8 AA.
 AC ABG93028;
 XX
 DT 20-NOV-2002 (first entry)
 XX
 DE Mouse class I MHC molecule Kb binding ovalbumin epitope.
 XX
 KW Regulator; transcription; cell death; phenotype; molecular scaffold;
 KW gene therapy; cancer; cardiovascular disease; arrhythmia; heart failure;
 KW ischaemia; obesity; neurodegenerative disease; Alzheimer's disease;
 KW bone pathology; dermatologic disease; psoriasis; infection; AIDS;
 KW acquired immunodeficiency syndrome; cosmetic; wound healing;
 KW antibiotic transport; drug toxicity; drug resistance; immunobiology;
 KW inflammation; allergic response; human immunodeficiency virus.
 XX
 OS Unidentified.
 XX
 FN WO2000262822-A2.
 XX

PD 15-AUG-2002.
 XX
 PF 04-FEB-2002; 2002WO-US02814.
 XX
 PR 02-FEB-2001; 2001US-265586P.
 PR 05-FEB-2001; 2001US-265860P.
 PR 27-FEB-2001; 2001US-271423P.
 PR 23-JAN-2001; 2001US-263226P.
 PR 28-MAR-2000; 2000US-192586P.
 PR 22-SEP-1997; 97US-935377P.
 XX
 PA (UVRP) UNIV ROCHESTER.
 XX
 PI Zauderer M, Smith ES;
 XX
 DR WPI; 2002-643398/69.
 XX
 PT Identifying regulator polypeptides which influence target
 PT transcriptional regulatory regions, useful for treating cancer,
 PT comprises introducing host cells expressing the polypeptide into a
 PT library of polynucleotides -
 XX
 PS Example 1; Page 82; 224pp; English.
 XX
 CC The invention discloses a method for identifying polynucleotides encoding
 CC a regulator polypeptide, whose expression induces activation of a target
 CC transcriptional regulatory region in a host cell. The method comprises
 CC providing a population of eukaryotic host cells capable of expressing the
 CC polypeptide, introducing into the host cell a library of polynucleotides
 CC encoding the polypeptides, permitting expression of the polypeptides and
 CC then recovering them from the host cells. The target transcriptional
 CC regulatory region is operably associated with a polynucleotide encoding a
 CC gene product, the expression of which results in host cell death or cause
 CC the host cells to exhibit a pre-determined modified phenotype and where
 CC the gene product is expressed upon activation of target transcriptional
 CC regulatory region. Each candidate regulator polypeptide comprises a
 CC candidate peptide and a molecular scaffold fused to the peptide so that
 CC the peptide is displayed on the surface of the candidate regulator
 CC polypeptide. The methods are useful in selecting and/or screening
 CC regulator molecules, such as polypeptides, which directly or indirectly
 CC induce or suppress the transcriptional activation of a target
 CC transcriptional regulatory region in a eukaryotic host cell. These
 CC regulator molecules may be used (e.g. in gene therapy) for preventing or
 CC treating cancers (e.g. breast or ovarian cancer), cardiovascular diseases
 CC (e.g. arhythmia, heart failure, ischaemia), obesity, neurodegenerative
 CC diseases (e.g. Alzheimer's disease), bone pathologies, dermatologic
 CC diseases (e.g. psoriasis), infections (e.g. viral, bacterial), acquired
 CC immunodeficiency syndrome (AIDS), in cosmetic applications and in wound
 CC healing. The method is also useful in screening regulator molecules that
 CC block antibiotic transport mechanisms, in drug toxicities and drug
 CC resistance applications and in improving the performance of existing or
 CC developmental drugs. It may also be used in immunobiology, inflammation,
 CC allergic response and in biotechnology applications. The sequences
 CC presented in ABG92946-ABG93029 are examples of regulator polypeptides.
 CC
 XX
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 23; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Db 1 STINFEKL 8
 1 STINFEKL 8
 XX
 AC AAE25400;
 XX
 ID AAE25400 standard; peptide; 8 AA.
 XX
 DT 30-OCT-2002 (first entry)

XX
 DE Ovalbumin peptide used in the invention.
 XX
 KW Recombinant vector; coat protein; CP; viral replication; infection;
 KW Zucchini yellow mosaic potyvirus; ZYMV; cucurbit fruit; vaccination;
 KW pharmaceutical; diagnostic; ovalbumin.
 XX
 OS Unidentified.
 XX
 PN WO200244323-A2.
 XX
 PD 06-JUN-2002.
 XX
 PF 28-NOV-2001; 2001WO-IL01098.
 XX
 PR 28-NOV-2000; 2000US-253136P.
 PR 27-SEP-2001; 2001US-0963761.
 XX
 PA (VIRO-) VIROGENE LTD.
 XX
 PI Gal-On A, Shioleth YM, Arazzi T, Ilan Y;
 XX
 DR WPI; 2002-537446/57.
 DR N-PSDB; AAD41429.
 XX
 PT Novel recombinant vector useful for transiently expressing heterologous
 PT peptide in plant comprises potyvirus nucleic acid sequence and
 PT heterologous sequence inserted at amino terminus of potyvirus coat
 PT protein -
 XX
 PS Claim 17; Page 60; 61pp; English.
 XX
 CC The invention relates to a recombinant vector for expressing a
 CC heterologous peptide at the amino-terminus of a potyvirus coat protein
 CC (CP). The vector includes sufficient potyvirus nucleic acid sequence
 CC to permit viral replication and spread within a plant infected by the
 CC vector. The invention also relates to Zucchini yellow mosaic potyvirus
 CC (ZYMV) Agri strain CP and its corresponding nucleic acid sequence. The
 CC recombinant vector is useful for transiently expressing a portion of
 CC the heterologous peptide in a plant. It is also useful for infecting a
 CC cucurbit fruit, is useful as a source of material for vaccination, a
 CC pharmaceutical or diagnostic application. The present sequence is a
 CC ovalbumin peptide used to fuse to the N-terminus of ZYMV Agri
 CC strain CP.
 XX
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 23; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Db 1 STINFEKL 8
 1 STINFEKL 8
 XX
 AC ABG31661;
 XX
 ID ABG31661 standard; Peptide; 8 AA.
 XX
 DT 05-NOV-2002 (first entry)
 XX
 DE Chicken ovalbumin MHC class I restricted epitope.
 XX
 KW Chicken; ovalbumin; MHC class I; major histocompatibility complex;
 KW polycationic compound; allergen; cytokine; chemokine; wound healing;
 KW cytotoxic drug; anti-oligogenic drug; immunostimulant; antiallergic;
 KW cytostatic; valine; immunogenic.
 XX
 OS Gallus gallus.
 XX

```

PN WO200253184-A2.
XX
XX 11-JUL-2002.
XX
XX 07-JAN-2002; 2002WO-EP00062.
PF
XX 05-JAN-2001; 2001WO-EP00087.
PR
XX 25-APR-2001; 2001AT-0000672.
XX
XX (INTE-) INTERCELL BIOMEDIZINISCHE FORSCHUNGS.
PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
XX
XX Lingnau K, Maltner F, Schmidt W, Birmstiel M, Buschle M;
XX
XX WPI; 2002-627324/67.
XX
XX Use of a polycationic compound for the preparation of a medicament with
XX retarded in vivo release
XX
XX Example 1; Page 13; 29pp; English.
XX
XX The invention relates to preparation of a medicament with retarded in
XX vivo release comprising use of a polycationic compound. The compound is
XX used in the preparation of a medicament with retarded in vivo release,
XX and a vaccine containing an antigen. The medicament includes e.g. an
XX allergen, a cytokine, a chemokine, an immunostimulatory nucleic acid, a
XX cytotoxic or an anti-oligogenic drug or a compound needed for wound
XX healing. The medicament prevents or ameliorates side effects of drugs,
XX which are due to its too fast distribution of the drug throughout the
XX body by exhibiting a retarded release of the drug from the site of
XX administration. In the case of vaccine the compounds provide a depot,
XX which allows a long lasting continuous and effective presentation of the
XX antigen to the immune system to create a protective immunity. This
XX sequence represents a chicken ovalbumin MHC class I restricted epitope
XX used in the scope of the invention.
XX
XX Sequence 8 AA;
XX
XX Query Match 100.0%; Score 38; DB 23; Length 8;
XX Best Local Similarity 100.0%; Pred. No. 9.3e+05;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
OY 1 SIINFEKL 8
DB 1 SIINFEKL 8
XX
XX RESULT 49
XX ABG31967
XX ID ABG31967 standard; Peptide; 8 AA.
XX
XX AC ABG31967;
XX
XX DT 05-NOV-2002 (first entry)
XX
XX DE Chicken ovalbumin OVA257-264-peptide.
XX
XX KW Chicken; ovalbumin; polycationic; antiinflammatory; immunostimulant;
XX anti-allergic; cycostatic; vulnerrary; medicament; inflammatory potential;
XX inflammation; vaccine; antigen; adjuvant; allergen; cytokine; chemokine;
XX immunostimulatory nucleic acid; cytotoxic drug; antioligogenic drug;
XX wound healing; OVA-peptide; epitope; major histocompatibility complex;
XX MHC.
XX
XX OS Gallus gallus.
XX
XX PN WO200253185-A2.
XX
XX PD 11-JUL-2002.
XX
XX PF 07-JAN-2002; 2002WO-EP00071.
XX
XX PR 05-JAN-2001; 2001WO-EP00087.
XX

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PR 25-APR-2001; 2001AT-0000670.
XX
XX (INTE-) INTERCELL BIOMEDIZINISCHE FORSCHUNGS.
PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
XX
XX Lingnau K, Eyed A, Schmidt W, Buschle M, Grill S;
XX
XX WPI; 2002-627325/67.
XX
XX Use of a polycationic compound for the preparation of a medicament with
XX reduced inflammatory potential
XX
XX Example 1; Page 13; 43pp; English.
XX
XX The invention discloses the use of a polycationic compound for the
XX preparation of a medicament with reduced inflammatory potential, for
XX treating or preventing inflammation or for a vaccine containing an
XX antigen, possible acting as an adjuvant. The medicaments include
XX allergens, cytokines, chemokines, immunostimulatory nucleic acids,
XX cytotoxic or antioligogenic drugs and compounds needed for wound healing.
XX The medicament acts locally at the site of administration, and lowers or
XX completely eliminates inflammatory side effects of medicaments. Thus the
XX medicament reduces the inflammatory potential of a medicament and allows
XX the administration of medicaments that are usually not administered or
XX only rarely administered due to their inflammatory side effects. The
XX sequence presented is the OVA 257-264-peptide, a major histocompatibility
XX complex (MHC) class I (H-2Kb) restricted epitope of chicken ovalbumin
XX which was used in the scope of the invention.
XX
XX Sequence 8 AA;
XX
XX Query Match 100.0%; Score 38; DB 23; Length 8;
XX Best Local Similarity 100.0%; Pred. No. 9.3e+05;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
OY 1 SIINFEKL 8
DB 1 SIINFEKL 8
XX
XX RESULT 50
XX AAU99718
XX ID AAU99718 standard; Peptide; 8 AA.
XX
XX AC AAU99718;
XX
XX DT 07-OCT-2002 (first entry)
XX
XX DE Mouse MHC class I Kb OVA peptide sequence.
XX
XX KW Mutant major histocompatibility complex class I chimeric protein; MHC;
XX lymphocyte; T-cell receptor; tissue sample; biopsy material; pathogen;
XX bodily fluid; T lymphocyte; neoplastic cell; tumour cell; MHC antigen;
XX virus; protozoan; bacteria; fungi; nematode; immune response; activator;
XX enhancer; T cell activator; mouse; recombinant yeast cell; Kb; OVA; Id;
XX beta2m; dev8.
XX
XX OS Mus sp.
XX
XX PN WO200246399-A2.
XX
XX PD 13-JUN-2002.
XX
XX PF 10-DEC-2001; 2001WO-US47817.
XX
XX PR 08-DEC-2000; 2000US-254495P.
XX
XX PA (UNII ) UNIV ILLINOIS FOUND.
XX
XX PI Kranz DM, Brophy S;
XX
XX DR WPI; 2002-527916/56.
XX

```

PT New isolated mutant major histocompatibility complex class I chimeric
 PT protein displayed on surfaces of recombinant yeast cells, has improved
 PT stability, and is useful for activating immune response -
 XX
 XX
 PS Example 3; Figure 18; 96pp; English.
 XX
 CC The present invention relates to a new mutant major histocompatibility
 CC complex (MHC) class I chimeric protein. The protein of the invention
 CC comprises a portion mediating binding to surfaces of recombinant yeast
 CC cells and a portion comprising peptide binding region of MHC class I
 CC protein, where the invention is improved in stability as compared with
 CC MHC class I chimeric protein which is not a mutant chimeric protein.
 CC The protein, further comprising a detectable label, is useful for
 CC detecting a lymphocyte having a T-cell receptor protein in a biological
 CC sample such as cells, tissue sample, biopsy material or bodily fluids.
 CC The method is useful for detecting a T lymphocyte that is specific for
 CC a neoplastic cell, a tumour cell, a virus-infected cell, a protozoan-
 CC infected cell, a bacterium-infected cell or a fungus-infected cell. The
 CC protein of the invention can be used to directly activate T cells, in
 CC order to identify/screen for peptide-MHC antigens. The protein is also
 CC useful in activating T cells that participate in the removal of target
 CC cells including neoplastic cells and cells infected with pathogenic
 CC agents including viruses, protozoans, bacteria, fungi or nematodes.
 CC The invention is improved in stability as compared with MHC class I
 CC protein which is not a mutant chimeric protein. The present amino acid
 CC sequence represents a mouse MHC peptide of the invention, as described
 CC above.
 XX
 SQ Sequence 8 AA;
 XX
 QY 1 SIINFEKL 8
 DB 1 SIINFEKL 8
 XX
 RESULT 51
 ABB08108
 ID ABB08108 standard; peptide; 8 AA.
 XX
 AC ABB08108;
 XX
 DT 10-SEP-2002 (first entry)
 XX
 DE Chicken ovalbumin CTL epitope fragment.
 XX
 KW T cell; antigen; tumour; vaccine; cytostatic; cancer; ovalbumin;
 KM CTL epitope.
 XX
 OS Gallus sp.
 OS
 PN US6387701-B1.
 XX
 PD 14-MAY-2002.
 XX
 PF 30-APR-1999; 99US-0302329.
 XX
 PR 30-APR-1996; 96US-0640444.
 PR 30-APR-1997; 97WO-US07317.
 PR 06-MAY-1998; 98US-0073819.
 PR 16-FEB-1999; 99US-0171916.
 XX
 PA (UYDU-) UNIV DUKE.
 PI Nair SK, Boczkowski DJ, Gilboa E;
 XX
 DR MPI; 2002-478447/51.
 XX
 PT Identifying tumor antigens that elicit T cell responses and which may
 PT be used for vaccinating against cancers, e.g. melanomas, breast

PT cancers, prostate cancers, colon cancers, and ovarian cancers,
 PT comprises a cytotoxicity assay -
 XX
 XX
 PS Example 1; Column 12; 21pp; English.
 XX
 CC The invention relates to identifying a tumour antigen that elicits a T
 CC cell response directed against the tumour. The antigen may then be used
 CC to vaccinate against cancers e.g. melanomas, bladder cancers, breast
 CC cancers, pancreatic cancers. The present sequence represents a chicken
 CC ovalbumin CTL epitope fragment.
 XX
 SQ Sequence 8 AA;
 XX
 QY 1 SIINFEKL 8
 DB 1 SIINFEKL 8
 XX
 RESULT 52
 ABB81273
 ID ABB81273 standard; peptide; 8 AA.
 XX
 AC ABB81273;
 XX
 DT 20-AUG-2002 (first entry)
 XX
 DE Chicken OVA 257-264 peptide SEQ ID NO:1.
 XX
 KW Yeast; dendritic cell; vaccine; immune response; ovalbumin; antifungal;
 KW immunostimulant; antibacterial; virucide; antiprotozoal; cytostatic;
 KW immunisation; cell mediated immunity; infectious disease; cancer.
 XX
 OS Gallus gallus.
 OS
 PN WO200239951-A2.
 XX
 PD 23-MAY-2002.
 XX
 PF 15-NOV-2001; 2001WO-US43537.
 XX
 PR 15-NOV-2000; 2000US-249173P.
 XX
 PA (GLOB-) GLOBE IMMUNE INC.
 PA (UYTE-) UNIV TECHNOLOGY CORP.
 XX
 PI Duke RC, Bellgrau D, Franzusoff A, Wilson CC;
 XX
 DR MPI; 2002-479895/51.
 XX
 PT Therapeutic composition, useful as vaccine, comprises dendritic cell
 PT intracellularly loaded with yeast vehicle and at least one antigen -
 XX
 PS Example 4; Page 34; 68pp; English.
 XX
 CC The present invention describes a therapeutic composition (I) comprising
 CC a dendritic cell (II), a yeast vehicle (III) and at least one antigen
 CC (IV), where (II) has been loaded intracellularly with (III) and (IV).
 CC (I) has immunostimulant, antibacterial, antifungal, virucide,
 CC antiprotozoal and cytostatic activities. (I) has many attributes that
 CC make it an ideal vaccine candidate, including ease of construction, low
 CC expense of mass production, biological stability and safety. No grossly
 CC adverse side effects of immunisation with whole yeast were apparent at
 CC the time of the initial vaccination or upon real administration. The
 CC composition provides a powerful strategy for the induction of cell-
 CC mediated immunity directed against a variety of infectious diseases and
 CC cancer targets. The present sequence represents a chicken OVA (ovalbumin)
 CC 257-264 peptide which is used in an example from the present invention.
 XX
 SQ Sequence 8 AA;
 XX

Query Match 100.0%; Score 38; DB 23; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 DB 1 SIINFEKL 8

RESULT 53

AAE22531
 ID AAE22531 standard; peptide; 8 AA.

AC AAE22531;

DT 26-JUL-2002 (first entry)

DE Ovalbumin peptide to evaluate the ability of aroC/siFA (P3H8) TML mutant.

KW SIFA: attenuated microorganism; medicament; allergen; gene therapy;

KW vaccine; protein therapy; virucide; hepatotropic; antiinflammatory;

KW protozoocide; ovalbumin.

OS Unidentified.

PN WO200226251-A1.

PD 04-APR-2002.

PF 01-OCT-2001; 2001WO-GB04358.

PR 29-SEP-2000; 2000GB-0023906.

PR 14-AUG-2001; 2001GB-0019802.

PA (MICR-) MICROSCIENCE LTD.

PI Brennan FR, Dougan G;

PS WPI; 2002-33986/37.

PT An attenuated Salmonella strain, for producing an elevated immune

PT response treat diseases, comprises a mutation that disrupts expression

PT of the sifa gene, and expresses a therapeutic heterologous peptide such

PT as an antigen -

PT Example 1; Page 7; 24pp; English.

PS The present invention relates to an attenuated Salmonella microorganism

PS which comprises a mutation that disrupts the expression of the sifa gene

PS and expresses a therapeutic heterologous peptide such as an antigen. The

PS attenuated microorganisms are useful for manufacturing medicaments to

PS treat or prevent a disease which can be treated by the heterologous

PS product and to increase the MHC class I-restricted response in a patient,

PS to deliver a therapeutic polynucleotide to a host cell to treat a disease

PS which can be corrected by administering the polynucleotide and to cause

PS an increase in the MHC class I-restricted response in a patient. They

PS are also useful to deliver heterologous antigens and allergens to a

PS patient, such as hepatitis, herpes simplex and Malarial antigens. The

PS method of the invention is useful for delivery of antisense nucleotides

PS or ribozymes in gene therapy. Sequences of the invention are also used

PS as vaccines and in protein therapy. The present sequence is ovalbumin

PS peptide (OVA 257-264) used to evaluate the ability of an aroC/siFA (P3H8)

PS TML mutant to stimulate a MHC class I-restricted response to heterologous

PS antigen.

SO Sequence 8 AA;

Query Match 100.0%; Score 38; DB 23; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8

DB 1 SIINFEKL 8

RESULT 54

ABB76050
 ID ABB76050 standard; peptide; 8 AA.

AC ABB76050;

DT 12-JUL-2002 (first entry)

DE Ovalbumin, H-2kb restricted epitope.

KW Bordetella pertussis; adenylcyclase; CyA; adenylcyclase;

KW vector; drug delivery; antigen delivery; cell targeting; CD11b;

KW ovalbumin; epitope; chicken.

OS Gallus sp.

PN EP1188446-A1.

PD 20-MAR-2002.

PF 15-SEP-2000; 2000EP-0402562.

PR 15-SEP-2000; 2000EP-0402562.

PA (INSP) INST PASTEUR.

PA (CNRS) CENT NAT RECH SCI.

PI Lelerc C, Guernonprez P, Ladant D, Guiso N, Khelif N;

PS WPI. 2002-354020/39.

PT Use of Bordetella adenylcyclase to make proteinaceous vector, useful

PT for drug or antigen delivery, selectively targets cells that express

PT CD11b -

PS Example B; Page 14; 34pp; English.

PS The present sequence is the peptide sequence of a chicken ovalbumin,

PS H-2kb restricted epitope, which was used as an experimental model

PS epitope in an example from the invention. The epitope was

PS genetically inserted into the catalytic domain of a detoxified, but

PS still invasive, mutant adenylate cyclase (adenylcyclase, CyA) of

PS Bordetella pertussis. The recombinant toxin, CyAOVA, was used to

PS immunise C57BL/6(H-2b) mice once i.v. CD4- and CD40-independent

PS cytotoxic T lymphocyte (CTL) priming was observed in the absence

PS of adjuvant. The invention relates to the novel use of Bordetella

PS CyA as a proteinaceous vector for targeting a molecule of interest

PS to the surface CD11b-expressing cells, especially dendritic cells

PS and neutrophils. The molecule of interest is translocated in the

PS cytosol to prime a CTL response. In a preferred embodiment, a

PS peptide is inserted into the catalytic domain of CyA at a

PS penultimate site. The peptide may be an intracellular bacterial

PS cell, tumour, viral, fungal or parasite cell antigen (all claimed).

PS Alternatively, a drug, especially an antiinflammatory, is chemically

PS coupled to CyA for drug delivery.

SO Sequence 8 AA;

Query Match 100.0%; Score 38; DB 23; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8

DB 1 SIINFEKL 8

RESULT 55

AAE19945

```

ID  AAE19945 standard; peptide; 8 AA.
XX
XX  AAE19945;
AC
XX  18-JUN-2002 (first entry)
DT
XX  Cytotoxic T-cell epitope for ovalbumin.
DE
XX  Cytotoxic T-cell; CTL; tumour; cancer; infection; cell-mediated immunity;
KW  vaccine; immune response; cytotoxic; T-cell epitope; ovalbumin.
KM
XX  Unidentified.
OS
XX  US2002018785-A1.
PN
XX  14-FEB-2002.
PD
XX  02-APR-2001; 2001US-0822250.
PF
XX  22-SEP-1997; 97US-0935377.
PR
XX  (UVRP ) UNIV ROCHESTER.
PA
XX  Zauderer M;
PI
XX
XX  WPI: 2002-239252/29.
DR
XX
XX  Representational Difference Analysis method for identification of
PT  antigens recognized by cytotoxic T cells and specific for human tumors,
PT  comprises improved selection of genes encoding target antigens -
XX
XX  Example 1; Page 13; 54pp; English.
PS
XX
XX  The present invention relates to novel methods for the identification
CC  of antigens recognised by cytotoxic T cells (CTLs) and specific for
CC  human tumors, cancers and infected cells. The method involves screening
CC  the products of an expression library generated from DNA/RNA of a cell
CC  expressing a target epitope with cytotoxic T cells generated against
CC  the cell to identify DNA clones expressing target epitope or providing
CC  cytotoxic T cells specific for a gene product differentially expressed
CC  by a cell and measuring the cross-reactivity of the cytotoxic T cells
CC  for cells expressing a target epitope in which the target epitope is
CC  identified as a gene product inducing cytotoxic T cells. The method is
CC  useful for identifying a target epitope or antigen specific for a tumour
CC  cell. The target epitope is also useful for identifying target antigens
CC  in other target cells against which it is desirable to induce cell-
CC  mediated immunity. The antigen identified by the method is useful
CC  in immunogenic compositions and vaccine preparations to induce the
CC  regression of tumours, cancers and infections in mammals. The invention
CC  also relates to vaccinia viral vectors which are useful for treating
CC  tumour-bearing mammals, including humans to generate immune response
CC  against the tumour cells. They are also useful for immunising or
CC  vaccinating tumour-free subjects to prevent tumour formation. The
CC  present sequence is cytotoxic T-cell epitope for ovalbumin. This
CC  peptide is used in the exemplification of the invention.
XX
SQ  Sequence 8 AA;

```

```

Query Match 100.0%; Score 38; DB 23; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 STINPEKL 8
   |||||
DB 1 STINPEKL 8

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RESULT 56
ABB09907
ID  ABB09907 standard; peptide; 8 AA.
XX
XX  ABB09907;
AC
XX

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DT 10-JUN-2002 (first entry)
XX
XX  Ovalbumin peptide (H-2kb CTL epitope).
DE
XX  Ovalbumin; cytokine; ELISPOT assay; polycationic substance;
KW  cytokine secreting cell; interleukin; IL; interferon; IFN; TNF; CSF;
KW  tumour necrosis factor; colony stimulating factor;
KW  enzyme-linked immunosorbent spot assay.
XX
XX  Aves.
OS
XX  WO200179854-A1.
PN
XX  25-OCT-2001.
PD
XX  12-APR-2001; 2001WO-EP04208.
PF
XX  13-APR-2000; 2000AT-0000645.
PR
XX  (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
PA
XX  Schallich J, Buschle M;
PI
XX
XX  WPI: 2002-240136/29.
DR
XX
XX  Identification of cytokine secreting cells involves incubating cell
PT  suspension containing antigen specific cells in the presence of
PT  polycationic substance -
XX
XX  Example 1; Page 10; 22pp; English.
PS
XX
XX  The sequence represents an ovalbumin peptide (H-2kb CTL epitope from
CC  ovalbumin). The invention relates to a novel method for identifying
CC  cytokine secreting cells (especially ELISPOT assays) using a polycationic
CC  substance. The method is useful for identifying cytokine secreting cells.
CC  The cytokine may be selected from interleukins (IL), interferons (IFN),
CC  tumour necrosis factors (TNF), and colony stimulating factors (CSF).
XX
XX
SQ  Sequence 8 AA;

```

```

Query Match 100.0%; Score 38; DB 23; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY 1 STINPEKL 8
   |||||
DB 1 STINPEKL 8

```

```

RESULT 57
AAU76942
ID  AAU76942 standard; Peptide; 8 AA.
XX
XX  AAU76942;
AC
XX
XX  05-JUN-2002 (first entry)
DT
XX
XX  OVA peptide (257-264).
DE
XX
XX  ODN; immunostimulatory oligodeoxynucleotide; Ovalbumin; OVA;
KW  carianat; vaccine; tuberculosis; diphteria; pertussis; measles;
KW  tetanus; acquired immune deficiency syndrome; AIDS; malaria;
KW  cardiovascular disease; cancer; deoxyinosine; chicken.
XX
XX  Gallus sp.
OS
XX  WO200193905-A1.
PN
XX  13-DEC-2001.
PD
XX  07-JUN-2001; 2001WO-EP06433.
PF
XX  08-JUN-2000; 2000AT-0001000.
PR

```


PR 23-NOV-2000; 2000AT-0001973.
 XX
 XX (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
 XX
 XX Schmidt W, Lingnau K, Schellack C, Egyed A;
 XX
 XX WPI; 2002-240927/29.
 DR
 XX
 XX
 PT New oligodeoxynucleic acid molecule useful as an immunostimulatory
 agent -
 XX
 XX Example 1; Page 18; 52pp; English.
 PS
 XX This invention relates to immunostimulatory oligodeoxynucleic acid
 CC molecules (ODN) that can be used to enhance an immune response for
 CC use in vaccines. The immunostimulatory oligonucleotides of the invention
 CC and pharmaceutical compounds containing them may be used as medicine,
 CC especially as an immunostimulatory agent, for the preparation of
 CC vaccines useful for the treatment of tuberculosis, diphtheria,
 CC pertussis, measles and tetanus, acquired immune deficiency syndrome
 CC (AIDS), malaria, cardiovascular diseases, and cancer. Oligonucleotides
 CC containing deoxyinosine residues (I-ODN) show a better immunostimulatory
 CC effect compared to prior art compounds containing Cpg motifs. The ODNs
 CC of the invention produce more specific immune response to a given
 CC antigen or antigen fragment than the prior art compounds containing Cpg.
 CC Using immunostimulatory oligonucleotides containing deoxyinosine reduces
 CC the induction of adverse side reactions, especially the induction of
 CC systemic TNF-alpha or interleukin-6. The immunostimulatory effect of the
 CC composition containing a polycationic polymer and an antigenic fragment
 CC was significantly higher than could be expected from the addition of the
 CC effects of each single component or even the addition of the effects of
 CC the ODN or the polycation with the antigen. The present sequence
 CC represents the ovalbumin (OVA) peptide used as an antigen in
 CC examples of the method of the invention.
 CC
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 23; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEKL 8
 |||||
 Db 1 SIINFEKL 8
 RESULT 58
 AAU76802
 ID AAU76802 standard; Peptide; 8 AA.
 AC
 XX AAU76802;
 XX
 DT 21-MAY-2002 (first entry)
 XX
 DE MHC class I-restricted epitope of chicken ovalbumin.
 XX
 XX Chicken; ovalbumin; immunostimulant; T cell epitope; inosine; cytosine;
 KM polycationic peptide; systemic immune response; MHC class I; vaccine;
 KM major histocompatibility complex class I.
 XX
 XX Gallus gallus.
 OS
 XX WO200193903-A1.
 PN
 XX 13-DEC-2001.
 PD
 PF 07-JUN-2001; 2001WO-EP06437.
 XX
 PR 08-JUN-2000; 2000AT-0001000.
 XX
 XX (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
 PA
 XX Egyed A, Lingnau K, Mattner F, Buschle M, Schmidt W;
 PI

XX
 DR WPI; 2002-205813/26.
 XX
 XX Pharmaceutical composition for the preparation of vaccine comprises T
 PT cell epitope(s) or its mixture, polycationic peptide and nucleic acid
 PT based on inosine and cytosine -
 XX
 XX Example 1; Page 10; 45pp; English.
 PS
 XX The invention relates to a pharmaceutical composition comprising a T cell
 CC epitope(s) or its mixture, a polycationic peptide and a nucleic acid
 CC based on inosine and cytosine. The composition of the invention induces a
 CC systemic immune response and is used for the preparation of a vaccine.
 CC This sequence represents an MHC class I-restricted epitope of chicken
 CC ovalbumin, used to test enhancement of immune response against an
 CC ovalbumin-derived peptide.
 CC
 SQ Sequence 8 AA;
 Query Match 100.0%; Score 38; DB 23; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEKL 8
 |||||
 Db 1 SIINFEKL 8
 RESULT 59
 AAU76869
 ID AAU76869 standard; Peptide; 8 AA.
 AC
 XX AAU76869;
 XX
 DT 21-MAY-2002 (first entry)
 XX
 DE OVA peptide fragment.
 XX
 KM Membrane vesicle; exosome; density cushion centrifugation;
 KM dendritic cell; MHC; major histocompatibility complex; CDI; tumour;
 KM immunotherapy treatment; cancer; infection; immune disease; antitumour;
 KM cytotatic; OVA.
 XX
 OS Unidentified.
 XX
 PN WO200182958-A2.
 XX
 PD 08-NOV-2001.
 XX
 PF 11-APR-2001; 2001WO-EP04173.
 XX
 PR 27-APR-2000; 2000US-0561205.
 PR 09-FEB-2001; 2001US-0780748.
 XX
 XX (APCE-) AP CELLS INC.
 PA
 XX Lamparski H, Ruegg C, Le Pecq J, Hsu D, Yao J;
 PI WPI; 2002-066489/09.
 DR
 XX Preparing membrane vesicle from biological sample for treating cancer,
 PT by culturing membrane vesicle-producing cells to release vesicles,
 PT enriching vesicles and subjecting sample to density cushion
 PT centrifugation -
 XX
 XX Example 12; Page 55; 103pp; English.
 PS
 XX The invention relates to a method for preparing membrane vesicles (in
 CC particular exosomes) from a biological sample, comprising culturing a
 CC population of membrane vesicle-producing cells under conditions allowing
 CC the release of the vesicles, enriching the vesicles and treating the
 CC enriched biological sample by density cushion centrifugation. Immunogenic
 CC membrane vesicles are useful for producing an immune response in a

CC subject, by obtaining a biological sample containing dendritic cells,
 CC isolating or purifying a membrane vesicle from the sample, contacting the
 CC purified vesicle with a peptide or a lipid under conditions allowing the
 CC peptide or lipid to bind an MHC or CD1 molecule at the surface of the
 CC vesicle, and administering the vesicle to the subject. Membrane vesicles
 CC are useful for immunotherapy treatment or prophylaxis of tumours, and for
 CC treating various disease conditions such as cancer, infections, and
 CC immune diseases. This sequence represents an OVA peptide fragment which
 CC is contacted with an exosome, to create an immunogenic membrane vesicle.

XX Sequence 8 AA;

Query Match 100.0%; Score 38; DB 23; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 |||||
 DB 1 SIINFEKL 8

RESULT 60

AAU11866
 ID AAU11866 standard; Peptide; 8 AA.

XX AAU11866;

DT 26-MAR-2002 (first entry)

DE Ovalbumin derived control peptide.

XX T0 terminator; PSA: DNA vaccine; anti-HIV, virucide;
 XX Human Immunodeficiency Virus; HIV; Gag; HIV gp120; HIV Pol; HIV Env;
 XX HIV VLP; measles fusion protein; measles haemagglutinin; epitope;
 XX measles nucleoprotein; influenza haemagglutinin; C3d gene; ovalbumin;
 XX cell-mediated immune response; humoral immune response; infection.

OS Unidentified.

PN WO200192470-A2.

PD 06-DEC-2001.

PF 02-MAR-2001; 2001WO-US06795.

PR 02-MAR-2000; 2000US-186364P.

PR 01-DEC-2000; 2000US-251083P.

PA (UYEM-) UNIV EMORY.

PI Robinson HL, Smith JM, Ross TM, Bright RA, Hua J, Ellenberger D;
 XX WPI; 2002-075465/10.

DR WPI; 2002-075465/10.

PT Novel pGA vector useful for immunising patient against measles,
 PT influenza has termination sequence encoding lambda T0 terminator and a
 PT eukaryotic transcription cassette with vaccine insert encoding
 PT immunogens of pathogens -
 XX
 XX Example 14; Page 61; 174pp; English.

CC The invention relates to a vector (a pGA construct) comprising a
 CC termination sequence coding for the lambda T0 terminator, a prokaryotic
 CC origin of replication, a selectable marker gene and a eukaryotic
 CC transcription cassette comprising a vaccine insert encoding one or more
 CC immunogens derived from a pathogen e.g. Human Immunodeficiency Virus
 CC (HIV) Gag, HIV gp120, HIV Pol, HIV Env, HIV VLP, or its mutants, measles
 CC fusion protein, measles haemagglutinin, measles nucleoprotein, influenza
 CC haemagglutinin, or its mutants, or subsequences, and optionally at least
 CC one C3d gene, is useful for immunising or treating a patient, when
 CC administered by an intramuscular or intradermal route. The immunisation
 CC methods using pGA elicit both cell-mediated and humoral immune responses
 CC that may limit the infection, spread or growth of the pathogen and result

CC in protection against subsequent challenge against the pathogen. The
 CC terminator sequence present prevents read-through from the kanamycin
 CC cassette into vaccine sequences while the plasmid is being produced in
 CC bacteria. Prevention of transcriptional read-through stabilises vaccine
 CC insert sequences by limiting the exposure of secondary structures that
 CC can be recognised by bacterial endonucleases. The present sequence
 CC is an ovalbumin control peptide used in an experiment to measure the
 CC T-cell response in monkeys inoculated with a pGA vector carrying vaccinia
 CC virus genes.

XX Sequence 8 AA;

Query Match 100.0%; Score 38; DB 23; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 |||||
 DB 1 SIINFEKL 8

RESULT 61

AAU11239
 ID AAU11239 standard; peptide; 8 AA.

XX AAU11239;

DT 12-MAR-2002 (first entry)

DE Immunodominant Kb-restricted Cytotoxic T lymphocyte epitope #2.

XX Cytostatic; vaccine; tetanus toxin; Frc; tumour; CTL;
 XX cytotoxic T-lymphocyte; immunodominant Kb-restricted CTL epitope.

OS Unidentified.

PN WO200179510-A1.

PD 25-OCT-2001.

PF 17-APR-2001; 2001WO-GB01719.

PR 17-APR-2000; 2000GB-0009470.

PA (CANC-) CANCER RES VENTURES LTD.

PI Rice J, Stevenson F;

XX WPI; 2002-066370/09.

PT Nucleic acid construct, useful to immunise against various diseases
 PT including cancer, expresses the first domain of tetanus toxin Frc fused
 PT to a disease peptide antigen to provide a vaccine -
 XX
 XX Disclosure; Page 25; 71pp; English.

CC The invention relates to a nucleic acid construct for delivery into
 CC living cells in vivo, to induce an immune response to a disease peptide
 CC antigen, where the construct directs expression of a fusion protein
 CC comprising the peptide antigen and the first domain of Frc. Also
 CC included are a nucleic acid vector comprising the above construct,
 CC a host cell comprising the above construct or vector and a method of
 CC producing a nucleic acid construct for inducing an immune response.
 CC The method comprises identifying a nucleic acid sequence encoding a
 CC disease peptide antigen comprising epitopes characteristic of the
 CC disease, cloning the nucleic acid sequence, introducing the cloned
 CC nucleic acid into a vector which allows the antigen to be expressed as a
 CC fusion with a first domain Frc from tetanus toxin, and optionally
 CC isolating the construct from the vector. The construct or vector is used
 CC as a vaccine to induce an immune response, particularly to tumour
 CC antigens. The present sequence is an immunodominant Kb-restricted
 CC cytotoxic T-lymphocyte (CTL) epitope suitable for inclusion in the
 CC vaccine of the invention.

XX Sequence 8 AA;
SQ

Query Match 100.0%; Score 38; DB 23; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 SIINFEKL 8
Db 1 SIINFEKL 8

RESULT 62
AAU09820
ID AAU09820 standard; peptide; 8 AA.

XX AAU09820;

DT 14-FEB-2002 (first entry)

DE Ovalbumin-derived class I H-2Kb restricted peptide (245).

KW Ovalbumin-derived class I H-2Kb restricted peptide (245); vaccine;
KW immunostimulant; immunogenic; interferon-gamma-producing T cell; IFN;
KW popliteal lymph node; spleen; immune response; systemic response.

OS Unidentified.

PN WO200178767-A2.

PD 25-OCT-2001.

PF 17-APR-2001; 2001WO-EP04313.

PR 14-APR-2000; 2000AT-0000657.

PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH.

PI Matner F, Zauner W, Schmidt W, Buschle M;

DR WPI; 2002-025970/03.

PT Pharmaceutical preparation for use as a potent vaccine for inducing an
PT improved immune response in a mammal, comprises a modified peptide -
PT

PS Example 1; Page 8; 18pp; English.

CC The invention relates to a pharmaceutical preparation comprising a
CC modified peptide, which induces an improved immune response in a mammal
CC compared to the wild type peptide. The neutral peptide (SIINFEKL) (one
CC negatively charged (Glu), one positively charged (Lys) amino acid) was
CC rendered negative by adding (at the N-terminus) Glu Glu or Glu Asp Glu
CC Asp, respectively. Results showed that the addition of 4 negatively-
CC charged amino acids (EDBD) at the N-terminus of peptide SIINFEKL makes
CC this peptide (in combination with poly-L-arginine) able to induce a high
CC amount of specific interferon (IFN)-gamma-producing T cells in the
CC draining (popliteal) lymph node (local response) and in the spleen
CC (systemic response). Thus, the addition of hydrophobic amino acids as
CC well as the addition of negatively charged amino acids transforms the
CC peptide SIINFEKL to a good inducer of specific T cells. The modified
CC peptides of the pharmaceutical composition induce a stronger immune
CC response in a mammal compared to wild type antigens. The present
CC sequence represents ovalbumin-derived class I H-2Kb restricted peptide
CC (245) used to produce the modified hydrophobic peptides described in
CC the method of the invention.

SQ Sequence 8 AA;

Query Match 100.0%; Score 38; DB 23; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 SIINFEKL 8

Db 1 SIINFEKL 8

RESULT 63
AAE13215
ID AAE13215 standard; peptide; 8 AA.

XX AAE13215;

DT 12-FEB-2002 (first entry)

DE Hen egg ovalbumin (OVA) peptide.

KW Cytotoxic T lymphocyte; CTL; T cell; tumour load; cancer radiotherapy;
KW immunostimulatory sequence oligonucleotide; ISS-ODN; chemotherapy;
KW immunosuppression; transplantation; autoimmune disease; infection;
KW acquired immune deficiency syndrome; AIDS; intracellular pathogen;
KW cytomegalovirus; mycobacterial infection; Epstein-Barr virus;
KW varicella zoster virus; human immunodeficiency virus; HIV; hen;
KW ovalbumin; OVA.

OS Unidentified.

PN WO200172123-A1.

PD 04-OCT-2001.

PF 28-MAR-2001; 2001WO-US10118.

PR 28-MAR-2000; 2000US-192537P.

PR 11-MAY-2000; 2000US-203567P.

PR 05-JUL-2000; 2000US-215895P.

PA (RECC) UNIV CALIFORNIA.

PA (VETE-) DEPT VETERANS AFFAIRS.

PI Raz E, Cho HJ, Richman DD, Horner AA;

DR WPI; 2002-010699/01.

PT Increasing antigen-specific cytotoxic T lymphocyte activity in a CD4+ T
PT cell deficient individual, useful to treat immunodeficiency and block
PT HIV infection, comprises administering immunostimulatory nucleic acid
PT

PS Example 1; Page 44; 91pp; English.

CC The present invention relates to a method for increasing antigen-specific
CC cytotoxic T lymphocyte (CTL) activity in a CD4+ T cell-deficient
CC individual, comprising administering an immunostimulatory sequence
CC oligonucleotide (ISS-ODN). The immunostimulatory nucleic acids of the
CC invention are used in CD4+ T cell-deficient individuals to decrease
CC tumour load, to treat a primary or acquired immunodeficiency,
CC particularly where the acquired immunodeficiency is temporary and due
CC to cancer radiotherapy or chemotherapy or immunosuppression following
CC bone marrow or organ transplantation, or autoimmune disease treatment,
CC or is acquired immunodeficiency syndrome (AIDS). The nucleic acids may
CC be used to treat a person at risk of becoming CD4+ T cell-deficient,
CC particularly where someone at risk of cancer recurrence. They are also
CC used to treat infection, particularly by an intracellular pathogen,
CC especially one caused by cytomegalovirus, Mycobacterium tuberculosis,
CC M. avium, Epstein-Barr virus, a fungus yeast, varicella zoster virus or
CC human immunodeficiency virus (HIV). The present sequence is a hen egg
CC ovalbumin (OVA) peptide, used in the exemplification of the invention.

SQ Sequence 8 AA;

Query Match 100.0%; Score 38; DB 23; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 SIINFEKL 8

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Db      1 STINFEKL 8
      |||||
RESULT 64
AAE13436
ID      AAE13436 standard; peptide; 8 AA.
XX
AC      AAE13436;
XX
DT      12-FEB-2002 (first entry)
XX
DE      Chicken ovalbumin major histocompatibility complex class I epitope.
XX
KW      Antigenic complex; epitope; heat shock protein; HSP; tether; javelin;
KW      major histocompatibility complex; MHC; therapy; immune response;
KW      malignancy; chicken.
XX
OS      Gallus gallus.
XX
PN      WO200179259-A1.
XX
PD      25-OCT-2001.
XX
PF      17-APR-2001; 2001WO-US12567.
XX
PR      17-APR-2000; 2000US-197462P.
XX
PA      (ROTH/) ROTHMAN J E.
PA      (MAYH/) MAYHEW M.
PA      (HOEW/) HOE M.
XX
PI      Rothman JE, Mayhew M, Hoe M;
XX
DR      WPI; 2002-017594/02.
XX
PT      A new antigenic complex comprising epitopes non-covalently joined to a
PT      heat shock protein by a molecular tether designated a javelin are
PT      useful to treat or prevent infectious disease or malignancy -
XX
XX      Example; Page 15; 47pp; English.
XX
PS      The present invention relates to an antigenic complex, comprising a
CC      number of epitopes non-covalently joined to a heat shock protein (HSP) by
CC      a tethering molecule referred to as javelin which has affinity for the
CC      HSP under physiological conditions, where the epitopes are covalently
CC      joined to the tethering molecule and one epitope is major
CC      histocompatibility complex class I (MHC) and the other MHC class II. The
CC      antigenic complex is used to induce immune responses directed towards the
CC      treatment or prevention of infectious diseases and malignancies. The
CC      present sequence is chicken ovalbumin MHC class I epitope.
XX
SQ      Sequence 8 AA;
XX
Query Match      100.0%; Score 38; DB 23; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY      1 STINFEKL 8
      |||||
DB      1 STINFEKL 8
XX
RESULT 65
AAU75056
ID      AAU75056 standard; peptide; 8 AA.
XX
AC      AAU75056;
XX
DT      23-APR-2002 (first entry)
XX
DE      Ovalbumin antigenic peptide.
XX

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KW      Ovalbumin; Th1; Th2; antigen; immunosuppressive; cytostatic;
KW      adjuvant; lipopolysaccharide; lipid; vaccine; immunogenicity;
KW      immunocompetence; autoimmune disease; infectious disease; OVA;
KW      graft-versus- host disease; tumour; transgenic T cell; chicken.
XX
OS      Gallus gallus.
XX
PN      WO200197838-A2.
XX
PD      27-DEC-2001.
XX
PF      18-JUN-2001; 2001WO-US19411.
XX
PR      16-JUN-2000; 2000US-212182P.
XX
PA      (BAYU ) BAYLOR RES INST.
XX
PI      Pulendran B, Bancheureau JF, Cutler CW;
XX
DR      WPI; 2002-114543/15.
XX
PT      Use of adjuvants comprising isolated lipid groups such as Porphyromonas
PT      gingivalis lipopolysaccharides or its detoxified forms or derivatives
PT      for preparation of compositions to elicit T-helper cell responses
PT      in mammals -
XX
PS      Disclosure; Page 20; 58pp; English.
XX
CC      This invention relates to the use of adjuvants comprising isolated lipid
CC      groups such as Porphyromonas gingivalis lipopolysaccharide, its
CC      detoxified forms or derivatives, for preparing compositions for
CC      eliciting Th2 responses, enhancing vaccine immunogenicity, modulating
CC      immunocompetence, treating autoimmune/infectious disease, stimulating
CC      interleukin-5 (IL-5)/IL-13 production or dampening interferon gamma
CC      production in mammal. The adjuvant or a pharmaceutical compound
CC      containing it is useful for enhancing antibody harvest in a laboratory
CC      animal through an elicited Th2 immune response, or by modulating Th2
CC      immune responses. The adjuvant is also useful to study the Th2
CC      immune response in laboratory animal research, for the treatment or
CC      prophylactic vaccination of humans or animals against graft-versus-
CC      host disease, and for treating tumours. The present sequence
CC      represents the Ovalbumin (OVA) peptide used as an antigen to
CC      stimulate transgenic T cells in the method of the invention.
XX
SQ      Sequence 8 AA;
XX
Query Match      100.0%; Score 38; DB 23; Length 8;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
OY      1 STINFEKL 8
      |||||
DB      1 STINFEKL 8
XX
RESULT 66
ABU07743
ID      ABU07743 standard; Peptide; 8 AA.
XX
AC      ABU07743;
XX
DT      23-MAY-2003 (first entry)
XX
DE      Chicken ovalbumin epitope presented on murine Kb MHC class I.
XX
KW      Chicken; ovalbumin; OVA; cytolysin; vaccine delivery; Kb MHC class I;
KW      intracellular delivery vehicle; nonvirulent bacterium; drug delivery;
KW      gene therapy; biosynthesis; high level protein delivery;
KW      major histocompatibility complex; cytosolic protein delivery.
XX
OS      Gallus gallus.
XX
PN      US2002142007-A1.
XX

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XX 03-OCT-2002.
PD 07-SEP-2001; 2001US-0949109.
XX
XX 13-AUG-1998; 98US-0133914.
PF 21-DEC-1999; 99US-0469197.
XX
XX (PORT/) PORTNOY D A.
PA (HIGGS/) HIGGINS D E.
XX
XX Portnoy DA, Higgins DE;
PI
XX WPI; 2003-328328/31.
DR
XX
XX New nonvirulent bacterium with genes coding for a non-secreted foreign
PT cytolysin or a different foreign agent, useful as an intracellular
PT delivery vehicle for delivering, e.g. vaccines, drugs or genes for
PT therapy to eukaryotic cells
XX
XX Example; Page 6; 14pp; English.
XX
XX The invention relates to a nonvirulent bacterium, which comprises a first
CC gene encoding a non-secreted foreign cytolysin operably linked to a
CC heterologous promoter and a second gene encoding a different foreign
CC agent. The nonvirulent bacterium is useful as an intracellular delivery
CC vehicle, particularly of agents to eukaryotic cells. The nonvirulent
CC bacterium is particularly useful for delivering foreign agents for
CC diagnosis, therapy (e.g. prophylaxis such as vaccine, delivery of
CC therapeutic drug, or gene therapy) or biosynthesis. The nonvirulent
CC bacterium is also useful for delivering nucleic acids that provide
CC templates for transcription or translation, or provide modulators of
CC transcription and/or translation. No protein purification is required
CC compared to prior art delivery systems. In addition, high levels of
CC protein can be delivered to the cytosol of virtually any cell and the
CC levels can be controlled through the use of inducible promoters. L.
CC monocytogenes LfO (listeriolysin or cytolysin) was transformed in
CC E.coli cells and used as a system to deliver chicken ovalbumin to the
CC cytosol of macrophages. The present sequence represents the chicken
CC ovalbumin epitope presented on murine Kb major histocompatibility
CC complex, MHC, class I.
XX
XX SQ Sequence 8 AA;
XX
XX Query Match
XX Best Local Similarity 100.0%; Score 38; DB 24; Length 8;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 SIINFEXL 8
XX |||||
XX Db 1 SIINFEXL 8
XX
XX RESULT 67
XX ABU08619
XX ID ABU08619 standard; Peptide; 8 AA.
XX
XX AC ABU08619;
XX
XX 23-MAY-2003 (first entry)
XX
XX DE Ovalbumin (OVA) residues 257-264.
XX
XX Cancer; tumour; antigen-presenting cell; APC; tumour cell conjugate;
XX cytokine; interleukin; interferon; IFN alpha; IFNbeta; IFNgamma;
XX tumour necrosis factor; TNF; transforming growth factor; TGF;
XX granulocyte-macrophage colony stimulating factor; GM-CSF; vaccine;
XX melanoma; kidney cancer; pulmonary carcinoma; hepatic carcinoma;
XX mammary cancer; prostatic carcinoma; gastric carcinoma; leukaemia;
XX ovalbumin-specific tetramer; (OVA)-specific tetramer; ovalbumin;
XX OVA.
XX
XX OS Gallus gallus.
XX
XX

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XX US2002182194-A1.
XX
XX 05-DEC-2002.
PD
XX
XX 25-MAR-2002; 2002US-0106173.
PF
XX
XX 04-APR-2001; 2001CN-0105852.
PR
XX
XX (SHAN-) SHANGHAI BRILLIANCE BIOTECH INST.
XX
XX Ju D, Tao Q, Ye D;
XX
XX WPI; 2003-328591/31.
DR
XX
XX New antigen-presenting cell and tumor cell conjugates, where the
PT antigen-presenting cell is modified by a cytokine gene, useful for the
PT preparation of a medicine for the therapy of cancer or a vaccine for
PT the prophylaxis of cancer
XX
XX Example 1; Page 4; 23pp; English.
XX
XX The invention describes an antigen-presenting cell (APC)/tumour cell
CC conjugate, where the APC is modified by a cytokine gene (interleukin
CC IL-2, IL-3, IL-4, IL-6, IL-12, IL-18, interferon (IFN) alpha, IFNbeta,
CC IFNgamma, tumour necrosis factor (TNF), transforming growth factor (TGF)
CC and/or granulocyte-macrophage colony stimulating factor (GM-CSF)). The
CC antigen-presenting cell/tumour cell conjugate is useful for the
CC preparation of a medicine for the therapy of cancer or a vaccine for the
CC prophylaxis of cancer. The cancer includes melanoma, kidney cancer,
CC pulmonary carcinoma, hepatic carcinoma, mammary cancer, prostatic
CC carcinoma, gastric carcinoma and leukaemia. This is the amino acid
CC sequence of Ovalbumin (OVA) residues 257-264 used in the creation of
CC Ovalbumin (OVA)-specific tetramers used in the vaccine of the invention.
XX
XX SQ Sequence 8 AA;
XX
XX Query Match
XX Best Local Similarity 100.0%; Score 38; DB 24; Length 8;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 SIINFEXL 8
XX |||||
XX Db 1 SIINFEXL 8
XX
XX RESULT 68
XX ABP57401
XX ID ABP57401 standard; peptide; 8 AA.
XX
XX AC ABP57401;
XX
XX 23-APR-2003 (first entry)
XX
XX DE Synthetic dimer peptide.
XX
XX Escherichia coli heat labile enterotoxin; Etx; cholera toxin; Ctx;
XX Vibrio cholerae; mutant; GM-1 ganglioside receptor; carrier; toxin;
XX virulence; cytostatic; vaccine; viral infection; cancer; EtxB; CtxB.
XX
XX OS Synthetic.
XX
XX WO2003000899-A1.
XX
XX PN 03-JAN-2003.
XX
XX PD 20-JUN-2002; 2002WO-GB02829.
XX
XX PR 22-JUN-2001; 2001GB-0015382.
XX
XX (UYBR-) UNIV BRISTOL.
XX
XX PA
XX
XX PI
XX
XX

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XX DR WPI; 2003-175291/17.
XX
XX PT Use of a mutant form of B subunit of Escherichia coli heat labile
XX PT enterotoxin or B subunit of cholera toxin for delivering an agent to a
XX PT target cell for treating viral infection or cancer
XX PS Example 5; Page 45; 84pp; English.
XX
XX CC The present invention describes a mutant form of B subunit of Escherichia
XX CC coli heat labile enterotoxin (EtxB) or B subunit of cholera toxin (CtxB)
XX CC from Vibrio cholerae which is useful for delivering an agent to a target
XX CC cell, and has GM-1 ganglioside receptor binding activity but has reduced
XX CC immunogenic and immunomodulatory activity relative to the wild-type form
XX CC of EtxB or CtxB. Also described: (1) treating a disease or condition in
XX CC a subject; (2) delivering the agent using the mutant to a target cell;
XX CC (3) a composition; and (4) a kit for delivering the agent to a target
XX CC cell. Mutant EtxB and CtxB have virulence and cytostatic activities and
XX CC can be used in vaccines. The mutant can be used for the preparation of
XX CC a medicament for delivering an exogenous peptide, which is the agent,
XX CC into the major histocompatibility complex (MHC) Class I antigen
XX CC processing and presenting pathways to elicit a cytotoxic T lymphocyte
XX CC (CTL) response, or for separate, simultaneous or combined use for
XX CC treating viral infection or cancer. The mutant form of EtxB or CtxB
XX CC enters mammalian cells without inducing a potent anti-B-subunit response
XX CC and immunomodulatory response. It may be linked with an agent to
XX CC upregulate the presentation of the antigen or antigenic determinant.
XX CC The present sequence represents a peptide which is used in an example
XX CC from the present invention.
XX
XX SQ Sequence 8 AA;
XX
XX Query Match 100.0%; Score 38; DB 24; Length 8;
XX Best Local Similarity 100.0%; Pred. No. 9.3e+05;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 SIINFEXL 8
XX |||||
XX 1 SIINFEXL 8
XX
XX DB
XX
XX RESULT 69
XX ABP58359
XX ID ABP58359 standard; Peptide; 8 AA.
XX
XX AC ABP58359;
XX
XX DT 07-APR-2003 (first entry)
XX
XX DE Ovalbumin-derived peptide OVA257-264.
XX
XX KW Chicken; ovalbumin; allergen; immunostimulant;
XX KW oligodeoxynucleic acid; ODN; vaccine.
XX
XX OS Gallus sp.
XX
XX PN WO200295027-A2.
XX
XX PD 28-NOV-2002.
XX
XX PF 17-MAY-2002; 2002WO-BE05448.
XX
XX PR 21-MAY-2001; 2001AT-0000805.
XX
XX PA (INTE-) INTERCELL BIOMEDIZINISCHE FORSCHUNGS.
XX PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
XX
XX PI Lingnan K, Schellack C, Schmidt W;
XX
XX DR WPI; 2003-183880/18.
XX
XX PT New oligodeoxynucleic acid molecules useful for the preparation of
XX PT vaccine -

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```

XX PS Example 3; Page 23; 57pp; English.
XX
XX CC The present sequence is that of OVA257-264, a major
XX CC histocompatibility complex class I (H-2Kb)-restricted epitope of
XX CC chicken ovalbumin. An example from the invention describes the
XX CC generation of specific immune responses against this allergen-derived
XX CC peptide using deoxyridine monophosphate-modified oligonucleotide
XX CC U-ODN 13 (see AB224776). U-ODN 13 is an example of new
XX CC oligodeoxynucleic acid (ODN) molecules useful in the preparation of
XX CC vaccines. The invention is based on the discovery that ODNs
XX CC containing deoxyridine residues have an immunostimulatory effect
XX CC comparable to, or greater than, ODNs containing CpG motifs.
XX CC Combining the ODN with an antigen strongly increases the potential
XX CC of the antigen to raise the protection/immune response of a
XX CC vaccinated individual.
XX
XX SQ Sequence 8 AA;
XX
XX Query Match 100.0%; Score 38; DB 24; Length 8;
XX Best Local Similarity 100.0%; Pred. No. 9.3e+05;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 SIINFEXL 8
XX |||||
XX 1 SIINFEXL 8
XX
XX DB
XX
XX RESULT 70
XX ABG73081
XX ID ABG73081 standard; Peptide; 8 AA.
XX
XX AC ABG73081;
XX
XX DT 02-APR-2003 (first entry)
XX
XX DE MHC Class I peptide OVA-8.
XX
XX KW Antigen-specific T lymphocyte; MHC-antigen complex; MHC Class I peptide;
XX KW major histocompatibility complex; tumour-specific killer T cell;
XX KW virus-specific killer T cell; cytostatic; virulence; OVA-8.
XX
XX OS Synthetic.
XX
XX PN US2002151690-A1.
XX
XX PD 17-OCT-2002.
XX
XX PF 05-NOV-1999; 99US-0434965.
XX
XX PR 12-AUG-1997; 97US-0909549.
XX
XX PA (LUXE/) LUXEMBURG A T.
XX PA (JACK/) JACKSON M R.
XX PA (PETE/) PETER P A.
XX
XX PI Luxembourg AT, Jackson MR, Peter PA;
XX
XX DR WPI; 2003-182532/18.
XX
XX PT Enriching antigen-specific T lymphocytes, for purifying or expanding in
XX PT vitro tumour- or virus-specific killer T cells for cell therapy,
XX PT comprises capture of the lymphocytes on a substrate coated with
XX PT antigenic peptide-MHC complexes -
XX
XX PS Example 2; Page 5; 40pp; English.
XX
XX CC The invention relates to a method for enriching antigen-specific T
XX CC lymphocytes, comprising contacting a heterogeneous population of
XX CC antigen-specific T lymphocytes with a matrix comprising MHC-antigen
XX CC complexes for a period of time sufficient to allow the antigen-specific T
XX CC lymphocytes to interact with the matrix, and eluting the antigen-specific
XX CC T lymphocytes from the matrix to provide an enriched population of

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CC antigen-specific T lymphocytes. The MHC-antigen complexes comprise one or
 CC more antigens. Also claimed is a matrix for capturing antigen-specific T
 CC lymphocytes, comprising a support having on its surface an immobilised
 CC Class I peptide and a predetermined amount of an antigen, or for
 CC capturing antigens, comprising a support having on its surface an
 CC immobilised empty Class I peptide which is capable of binding one or more
 CC antigens, and isolating antigen-specific T lymphocytes from a
 CC heterogeneous population of cells from a patient. The methods are useful
 CC for enriching antigen-specific T lymphocytes to purify and expand in
 CC vitro tumour and virus-specific killer T cells for cell therapy. The
 CC methods are also useful for isolating or preparing a population of
 CC antigen-specific T lymphocytes from a patient for treatment of the
 CC patient's disease or condition. This sequence represents an MHC Class I
 CC peptide used in the method of the invention.

CC Sequence 8 AA;

Query Match 100.0%; Score 38; DB 24; Length 8;
 Best Local Similarity 100.0%; Pred. No. 9.3e+05;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 |||||
 DB 1 SIINFEKL 8

RESULT 71

ABP56760
 ID ABP56760 standard; peptide; 8 AA.

AC ABP56760;

DT 31-MAR-2003 (first entry)

DE Ovalbumin derived peptide OVA 257-264.

XX Stabilisation; polycationic polymer; medicine; vaccination;

KW gene therapy; drug; ovalbumin.

OS Gallus sp.

OS Synthetic.

PN WO200294845-A2.

PD 28-NOV-2002.

PF 17-MAY-2002; 2002WO-EP05447.

PR 21-MAY-2001; 2001AT-0000805.

PA (INTE-) INTERCELL BIOMEDIZINISCHE FORSCHUNGS.

PI (CIST-) CISTEM BIOTECHNOLOGIES GMBH.

PI Schellack C, Lingnau K, Schmidt W;

XX WPI; 2003-140356/13.

DR Use of polycationic polymer for stabilisation of nucleic acids -

XX Example 1; Page 6; 28pp; English.

PS The present invention describes a method for the stabilisation of nucleic
 CC acids involving contacting nucleic acids with a polycationic polymer in
 CC aqueous solution or suspension. The method can be used for the
 CC stabilisation of nucleic acid which can be used in medicines e.g.
 CC vaccination, as a general immunostimulant, as an antiseptic drug
 CC or gene therapy drug. The present sequence represents an ovalbumin
 CC derived peptide, designated OVA 257-264, which is used in an example
 CC from the present invention.

XX Sequence 8 AA;

QY Query Match

100.0%; Score 38; DB 24; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 |||||
 DB 1 SIINFEKL 8

RESULT 72

ABP60027
 ID ABP60027 standard; Peptide; 8 AA.

AC ABP60027;

DT 07-MAR-2003 (first entry)

DE Ovalbumin antigenic peptide.

XX TOP; thimet oligopeptidase; EC3.4.25.15; cytostatic; tumour;

KW immunostimulant; major histocompatibility complex class I; MHC;

KW T-cell immunity; chicken.

OS Gallus gallus.

PN WO200279388-A2.

PD 10-OCT-2002.

PF 01-APR-2002; 2002WO-US10385.

PR 30-MAR-2001; 2001US-280669P.

PA (UTMA-) UNIV MASSACHUSETTS.

PI Rock KL, Goldberg AL;

XX WPI; 2003-103265/09.

DR New recombinant cell comprising an exogenously derived nucleic acid

PT coding for a thimet oligopeptidase polypeptide, useful for modulating

PT an antigenic response in a mammal for treating e.g., tumour -

XX Example 1; Page 50; 73pp; English.

PS The invention relates to a new recombinant cell comprising an exogenously

CC derived nucleic acid that codes for a thimet oligopeptidase (TOP)

CC polypeptide. The TOP polypeptide is overexpressed in the cell compared to

CC a wild-type cell from which the recombinant cell is derived. The activity

CC of TOP may be described as cytostatic and immunostimulatory. Thimet

CC oligopeptidase (TOP; EC3.4.25.15) plays a key role in modulating levels

CC of major histocompatibility complex (MHC) class I-presented peptides. The

CC recombinant host cell of the invention is useful for modulating an

CC antigenic response in a mammal. Methods of the invention are useful for

CC screening a test compound for its ability to serve as an immunomodulatory

CC agent and identifying an antigen resistant to thimet oligopeptidase

CC degradation. A method of the invention is useful for increasing CD8

CC T-cell immunity, which uses vaccination with a TOP inhibitor for

CC decreasing TOP expression or activity. The vaccination method uses

CC treated tumour cells, antigen bearing/pulsed dendritic cells or injection

CC of a viral vector. The recombinant host cell is useful for treating

CC tumours. The current sequence represents an ovalbumin antigenic peptide

CC that is used in an example from the invention.

XX Sequence 8 AA;

Query Match 100.0%; Score 38; DB 24; Length 8;

Best Local Similarity 100.0%; Pred. No. 9.3e+05;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 |||||
 DB 1 SIINFEKL 8

```

RESULT 73
ABU1029
ID ABU1029 standard; Peptide; 8 AA.
XX
XX ABU1029;
XX
DT 04-FEB-2003 (first entry)
XX
DE Ovalbumin immunodominant epitope.
XX
KW Epitope; antigen-specific immunoglobulin; Ig; early/late promoter;
KW heavy chain constant region; light chain constant region;
KW variable region; camelised Ig heavy chain variable region; MHC; CTL;
KW major histocompatibility class; cytotoxic T-lymphocyte.
XX
OS Unidentified.
XX
PN US2002123057-A1.
XX
PD 05-SEP-2002.
XX
PF 14-NOV-2001; 2001US-0987456.
XX
PR 17-NOV-2000; 2000US-249268P.
PR 18-JAN-2001; 2001US-262067P.
PR 27-FEB-2001; 2001US-271424P.
PR 15-JUN-2001; 2001US-298087P.
XX
PA (UYRP ) UNIV ROCHESTER.
XX
PI Zauderer M, Smith ES;
XX
DR WPI; 2003-066785/06.
XX
PT Selecting polynucleotides which encode antigen-specific immunoglobulin
PT molecules, by introducing the library of polynucleotides into the host
PT cells, and recovering the polynucleotides of the library for the
PT antigen -
XX
XX Example 5; Page 45; 108pp; English.
XX
XX The invention relates to selecting polynucleotides which encode antigen
XX -specific immunoglobulins (Ig) (or fragments) comprising introducing into
XX a population of host cells, a 1st and 2nd library of polynucleotides
XX encoding, several 1st and 2nd Ig subunit polypeptides, permitting
XX expression of Ig molecules (via control element e.g. an early/late
XX promoter), contacting Ig molecules with an antigen,
XX and recovering polynucleotides of the 1st library for the antigen.
XX The Ig molecules are heavy and light chain constant regions and
XX variable regions linked via peptide linkers and optionally directed via
XX signal peptides or transmembrane domains to different cell compartments.
XX Also included is a method of selecting polynucleotides which encode a
XX single-domain antigen-specific Ig molecule (its anti-specific fragment),
XX by: (a) introducing into a population of eukaryotic host cells capable of
XX expressing the Ig molecule a library of polynucleotides encoding
XX (through operable association with a transcriptional control region)
XX several single-domain Ig polypeptides (each comprising a Ig heavy chain
XX constant region, a camelised Ig heavy chain variable region, and a
XX signal peptide capable of directing cell surface expression or
XX secretion of Ig subunit polypeptide); (b) permitting expression of Ig
XX molecules (or antigen-specific fragments) from the host cells;
XX (c) contacting the Ig molecules with an antigen; and (d) recovering
XX polynucleotides of the library from those host cells expressing Ig
XX molecules which bind the antigens. The methods are useful for selecting
XX polynucleotides which encode an antigen-specific Ig molecule, or its
XX fragment. The present sequence is a major histocompatibility class II
XX (MHC II), cytotoxic T-lymphocyte (CTL) epitope expressed on the surface
XX of host cells used in the method of the invention.
XX
SQ Sequence 8 AA;
Query Match 100.0%; Score 38; DB 24; Length 8;

```

```

Beat Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
   |||||
Db 1 SIINFEKL 8

RESULT 74
AAB84323
ID AAB84323 standard; peptide; 9 AA.
XX
XX AAB84323;
XX
AC AAB84323;
XX
DT 22-AUG-2001 (first entry)
XX
DE Peptide used to produce IemA peptide variants.
XX
XX IemA; CD8+ epitope; T cell response.
XX
XX Synthetic.
XX
XX WO200140275-A2.
XX
PN 07-JUN-2001.
XX
PD 06-DEC-2000; 2000WO-US33027.
XX
PR 06-DEC-1999; 99US-0169227.
XX
PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
PI Kurlander RJ, Chao E, Fields J;
XX
DR WPI; 2001-389952/41.
XX
PT New isolated variant of IemA, IemA, comprising a hydrophobic element
PT joined to a CD8+ epitope, useful for inducing a directed CD8+ T cell
PT response or as a treatment or prophylactic against diseases -
XX
XX Disclosure; Page 7; 65pp; English.
XX
XX The specification describes a peptide variant of IemA, comprising a
XX hydrophobic element joined to a CD8+ epitope. The peptides may be
XX used therapeutically by administering the peptides to a patient having
XX a need to induce a directed CD8+ T cell response. The peptide may also
XX be used as a preventive measure to avoid a disease or condition, or to
XX treat subjects already afflicted with a disease. The present sequence
XX was used to create peptides of the invention.
XX
SQ Sequence 9 AA;
Query Match 100.0%; Score 38; DB 22; Length 9;
Best Local Similarity 100.0%; Pred. No. 9.3e+05;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
   |||||
Db 2 SIINFEKL 9

RESULT 75
ABP57402
ID ABP57402 standard; peptide; 9 AA.
XX
XX ABP57402;
XX
AC ABP57402;
XX
DT 23-APR-2003 (first entry)
XX
DE Synthetic 9mer peptide.
XX
XX Escherichia coli heat labile enterotoxin; Etx; cholera toxin; Ctx;
XX
XX

```


KM	Vibrio cholerae; mutant; GM-1 ganglioside receptor; carrier; toxin;
KW	vincic; cytostatic; vaccine; viral infection; cancer; EtxB, CtxB.
OS	Synthetic.
XX	
PN	WO200300899-A1.
PD	03-JAN-2003.
PP	20-JUN-2002; 2002MO-GB02829.
PR	22-JUN-2001; 2001GB-0015382.
PA	(UYBR-) UNIV BRISTOL.
B1	Hirst TR;
DR	WPI; 2003-175291/17.
PT	Use of a mutant form of B subunit of Escherichia coli heat labile enterotoxin or B subunit of cholera toxin for delivering an agent to a target cell for treating viral infection or cancer
PS	Example 5; Page 45; 84pp; English.
CC	The present invention describes a mutant form of B subunit of Escherichia coli heat labile enterotoxin (EtxB) or B subunit of cholera toxin (CtxB) from Vibrio cholerae which is useful for delivering an agent to a target cell, and has Gm-1 ganglioside receptor binding activity but has reduced immunogenic and immunomodulatory activity relative to the wild-type form of EtxB or CtxB. Also described: (1) treating a disease or condition in a subject; (2) delivering the agent using the mutant to a target cell; (3) a composition; and (4) a kit for delivering the agent to a target cell. Mutant EtxB and CtxB have virulence and cytopathic activities and can be used in vaccines. The mutant can be used for the preparation of a medicament for delivering an exogenous peptide, which is the agent, into the major histocompatibility complex (MHC) Class I antigen processing and presenting pathways to elicit a cytotoxic T lymphocyte (CTL) response, or for separate, simultaneous or combined use for treating viral infection or cancer. The mutant form of EtxB or CtxB enters mammalian cells without inducing a potent anti-B-subunit response and immunomodulatory response. It may be linked with an agent to upregulate the presentation of the antigen or antigenic determinant. The present sequence represents a peptide which is used in an example from the present invention.
SC	Sequence 9 AA;
OY	Query Match 100.0%; Score 38; DB 24; Length 9; Best Local Similarity 100.0%; Pred. No. 9.36+05; Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
DQ	1 SIINPEKL 8 2 SIINPEKL 9
ID	AA04643 standard; peptide; 10 AA.
AC	AA04643;
DT	01-AUG-1997 (first entry)
DE	Ovalbumin-derived activated CD8+ T cells epitope OVA10N.
KM	Macrophage; artificial antigen presenting cell; APC; cancer;
KW	tumour; neoplasia; viral infection; retroviral infection;
OS	autoimmune.
XX	
XX	Synthetic.

```

PN      MO9637107-A1.
XX
XX      28-NOV-1996.
XX
XX      22-MAY-1996;      96WO-US07436.
XX
XX      23-MAY-1995;      95US-044761.
XX
XX      (SCRI ) SCRIpps RES INST.
XX
XX      DebruJn MLH, Jackson MR, Peterson PA;
XX
XX      WPI; 1997-020850/02.
XX
XX      Prodn. of activated CD8+ T cells directed to specific antigen - can
PT      specifically kill target cells useful to treat, e.g. cancer
XX
XX      Example 1; Page 26; 84pp; English.
XX
XX      The method for the production of activated CD8+ T cells specifically
CC      directed towards a particular antigen involves affixing peptides
CC      corresponding to the particular antigen to an artificial support;
CC      contacting macrophages with the affixed peptides for a time sufficient
CC      for the peptides to be engulfed, and at least a portion of the peptides
CC      to be presented on the surface of the macrophage; and contacting
CC      unprimed CD8+ T cells with the peptide presenting macrophages for a
CC      time sufficient to activate the unprimed CD8+ T cells. The present
CC      sequence represents a peptide designated OVA10N which corresponds to
CC      ovalbumin, a Kb-restricted peptide antigen. This represents the optimal
CC      peptide with the addition of two amino acids at th amino-terminus.
CC      Small extensions to the optimal peptide affect the affinity of the
CC      peptide for soluble class I molecules in vitro e.g. the addition
CC      of two amino acids to the amino-terminus lowers the affinity to Kb by
CC      76-fold compared to the optimal peptide; addition of two amino acids to
CC      the carboxy-terminus lowers the affinity by 4-fold. The method,
CC      macrophages and artificial antigen presenting cell, having a peptide
CC      corresponding to the particular antigen present on its surface and at
CC      least a portion of an artificial support in its interior, can be used to
CC      treat conditions (e.g. cancer, tumours, neoplasia, viral or retroviral
CC      infection or autoimmune or autoimmune-type conditions) in patients via
CC      the specific killing of target cells.
XX
XX      Sequence      10 AA;
SQ
XX
XX      Query Match      100.0%; Score 38; DB 18; Length 10;
XX      Best Local Similarity 100.0%; Pred. No. 0.15;
XX      Matches      8; Conservative 0; Mismatches 0; Indels 0; Gaps 0,
XX
XX      1 SIINFEKL 8
XX      |||||
XX      3 SIINFEKL 10
XX
XX      RESULT 77
XX      ID      AAW04644 standard; peptide; 10 AA.
XX
XX      AAW04644;
XX
XX      01-AUG-1997 (first entry)
XX
XX      Ovalbumin-derived activated CD8+ T cells epitope OVA10C.
XX
XX      Macrophage; artificial antigen presenting cell; APC; cancer;
XX      tumour; neoplasia; viral infection; retroviral infection;
XX      autoimmune.
XX
XX      Synthetic.
XX
XX      MO9637107-A1.
XX
XX      28-NOV-1996.
XX

```

PF 22-MAY-1996; 96WO-US07436.
 XX
 PR 23-MAY-1995; 95US-0447761.
 XX
 PA (SCRI) SCRIPPS RES INST.
 XX
 PI DeBrujn MLH, Jackson MR, Peterson PA;
 XX
 DR WPI, 1997-020850/02.
 XX
 PT Prodn. of activated CD8+ T cells directed to specific antigen - can
 PT specifically kill target cells useful to treat, e.g. cancer
 XX
 PS Example 1; Page 26; 84pp; English.
 XX
 CC The method for the production of activated CD8+ T cells specifically
 CC directed towards a particular antigen involves affixing peptides
 CC corresponding to the particular antigen to an artificial support;
 CC contacting macrophages with the affixed peptides for a time sufficient
 CC for the peptides to be engulfed, and at least a portion of the peptides
 CC to be presented on the surface of the macrophage; and contacting
 CC unprimed CD8+ T cells with the peptide presenting macrophages for a
 CC time sufficient to activate the unprimed CD8+ T cells. The present
 CC sequence represents a peptide designated OVA10C which corresponds to
 CC ovalbumin, a Kb-restricted peptide antigen. This represents the optimal
 CC peptide with the addition of two amino acids at the carboxy-terminus.
 CC Small extensions to the optimal peptide affect the affinity of the
 CC peptide for soluble class I molecules in vitro e.g. the addition
 CC of two amino acids to the amino-terminus lowers the affinity to Kb by
 CC 76-fold compared to the optimal peptide; addition of two amino acids to
 CC the carboxy-terminus lowers the affinity by 4-fold. The method,
 CC macrophages and artificial antigen presenting cell, having a peptide
 CC corresponding to the particular antigen present on its surface and at
 CC least a portion of an artificial support in its interior, can be used to
 CC treat conditions (e.g. cancer, tumours, neoplasia, viral or retroviral
 CC infection or autoimmune or autoimmune-type conditions) in patients via
 CC the specific killing of target cells.
 XX
 SQ Sequence 10 AA;
 XX
 Query Match 100.0%; Score 38; DB 18; Length 10;
 Best Local Similarity 100.0%; Pred. No. 0.15;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEXL 8
 DB 1 SIINFEXL 8
 XX
 RESULT 78
 AAU09821
 ID AAU09821 standard; peptide; 10 AA.
 XX
 AC AAU09821;
 XX
 DT 14-FEB-2002 (first entry)
 XX
 DE Modified ovalbumin-derived class I H-2Kb restricted peptide #1.
 XX
 KW Ovalbumin-derived class I H-2Kb restricted peptide; vaccine;
 KW immunostimulant; immunogenic; interferon-gamma-producing T cell; IFN;
 KW popliteal lymph node; spleen; immune response; systemic response.
 XX
 OS Synthetic.
 OS
 PN WO200178767-A2.
 XX
 PD 25-OCT-2001.
 XX
 PF 17-APR-2001; 2001WO-EP04313.
 XX
 PR 14-APR-2000; 2000AT-0000657.
 XX

PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
 XX
 PI Mattner F, Zauner W, Schmidt W, Buschle M;
 XX
 DR WPI, 2002-025970/03.
 XX
 PT Pharmaceutical preparation for use as a potent vaccine for inducing an
 PT improved immune response in a mammal, comprises a modified peptide -
 XX
 PS Example 1; Page 9; 18pp; English.
 XX
 CC The invention relates to a pharmaceutical preparation comprising a
 CC modified peptide, which induces an improved immune response in a mammal
 CC compared to the wild type peptide. The neutral peptide (SIINFEXL) (one
 CC negatively charged (Glu), one positively charged (Lys) amino acid) was
 CC rendered negative by adding (at the N-terminus) Glu Glu or Glu Asp Glu
 CC Asp, respectively. Results showed that the addition of 4 negatively-
 CC charged amino acids (EDSD) at the N-terminus of peptide SIINFEXL makes
 CC this peptide (in combination with poly-L-arginine) able to induce a high
 CC amount of specific interferon (IFN)-gamma-producing T cells in the
 CC draining (popliteal) lymph node (local response) and in the spleen
 CC (systemic response). Thus, the addition of hydrophobic amino acids as
 CC well as the addition of negatively charged amino acids transforms the
 CC peptide SIINFEXL to a good inducer of specific T cells. The modified
 CC peptides of the pharmaceutical composition induce a stronger immune
 CC response in a mammal compared to wild type antigens. The present
 CC sequence represents modified ovalbumin-derived class I H-2Kb restricted
 CC peptide #1 as described in the method of the invention.
 XX
 SQ Sequence 10 AA;
 XX
 Query Match 100.0%; Score 38; DB 23; Length 10;
 Best Local Similarity 100.0%; Pred. No. 0.15;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEXL 8
 DB 3 SIINFEXL 10
 XX
 RESULT 79
 AAU09825
 ID AAU09825 standard; peptide; 10 AA.
 XX
 AC AAU09825;
 XX
 DT 14-FEB-2002 (first entry)
 XX
 DE Modified ovalbumin-derived class I H-2Kb restricted peptide #5.
 XX
 KW Ovalbumin-derived class I H-2Kb restricted peptide; vaccine;
 KW immunostimulant; immunogenic; interferon-gamma-producing T cell; IFN;
 KW popliteal lymph node; spleen; immune response; systemic response.
 XX
 OS Synthetic.
 OS
 PN WO200178767-A2.
 XX
 PD 25-OCT-2001.
 XX
 PF 17-APR-2001; 2001WO-EP04313.
 XX
 PR 14-APR-2000; 2000AT-0000657.
 XX
 PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
 XX
 PI Mattner F, Zauner W, Schmidt W, Buschle M;
 XX
 DR WPI, 2002-025970/03.
 XX
 PT Pharmaceutical preparation for use as a potent vaccine for inducing an
 PT improved immune response in a mammal, comprises a modified peptide -
 XX

PS Example 2; Page 9; 18pp; English.
 XX
 CC The invention relates to a pharmaceutical preparation comprising a
 CC modified peptide, which induces an improved immune response in a mammal
 CC compared to the wild type peptide. The neutral peptide (SIINFEKL) (one
 CC negatively charged (Glu), one positively charged (Lys) amino acid) was
 CC rendered negative by adding (at the N-terminus) Glu Glu or Glu Asp Glu
 CC Asp, respectively. Results showed that the addition of 4 negatively-
 CC charged amino acids (EDBD) at the N-terminus of peptide SIINFEKL makes
 CC this peptide (in combination with poly-L-arginine) able to induce a high
 CC amount of specific interferon (IFN)-gamma-producing T cells in the
 CC draining (popliteal) lymph node (local response) and in the spleen
 CC (systemic response). Thus, the addition of hydrophobic amino acids as
 CC well as the addition of negatively charged amino acids transforms the
 CC peptide SIINFEKL to a good inducer of specific T cells. The modified
 CC peptides of the pharmaceutical composition induce a stronger immune
 CC response in a mammal compared to wild type antigens. The present
 CC sequence represents modified ovalbumin-derived class I H-2Kb restricted
 CC peptide #5 as described in the method of the invention.
 SQ Sequence 10 AA;
 Query Match 100.0%; Score 38; DB 23; Length 10;
 Best Local Similarity 100.0%; Pred. No. 0.15;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 SIINFEKL 8
 |||||
 3 SIINFEKL 10
 Db
 RESULT 80
 AAW14122
 ID AAW14122 standard; peptide; 12 AA.
 AC AAW14122;
 XX
 DT 20-OCT-1997 (first entry)
 XX
 DE OVA protein derived MHC class I binding peptide.
 XX
 XX Major histocompatibility complex; MHC; target; binding; tumour;
 KW cancer; neoplasia; LSTRA; EL-4; identification; detection; screening;
 KW tissue typing; Bcr-abl; IFV; Influenza.
 XX
 OS Mus sp.
 XX
 XX Key Location/Qualifiers
 FH Misc-difference 12
 FT Misc-difference /note="bioinylated"
 FT
 XX
 PN WO9641188-A1.
 XX
 PD 19-DEC-1996.
 XX
 PF 07-JUN-1996; 96WO-US09680.
 XX
 PR 07-JUN-1995; 95US-0485610.
 XX
 PA (UNITW) UNIV WASHINGTON.
 XX
 PI Cheever MA, Chen W;
 XX
 DR WPI; 1997-108657/10.
 XX
 PT Identifying major histocompatibility complex class I binding mols. -
 PT using peptide(s) having a core of 7-14 amino acids with extra amino
 PT acids and a reporter gp. at the N- or C-terminus, useful for tissue
 PT typing
 XX
 PS Example 3; Page 23; 41pp; English.
 CC AAW14122 is a bioinylated peptides derived from the OVA (ovalbumin)

CC protein which can be obtained from either an LSTRA or EL-4 tumour
 CC of Balb/c mice. The peptides bind to MHC class I molecules. This
 CC is useful for tissue typing or for screening for molecules that
 CC interact with MHC class I molecules. MHC class I molecules can be
 CC identified using the peptides and also the peptides are useful in
 CC vaccines against disease and infection e.g. caused by viruses,
 CC bacteria or tumours.
 CC
 SQ Sequence 12 AA;
 Query Match 100.0%; Score 38; DB 18; Length 12;
 Best Local Similarity 100.0%; Pred. No. 0.18;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 SIINFEKL 8
 |||||
 1 SIINFEKL 8
 Db
 RESULT 81
 AAU09822
 ID AAU09822 standard; peptide; 12 AA.
 AC AAU09822;
 XX
 DT 14-FEB-2002 (first entry)
 XX
 DE Modified ovalbumin-derived class I H-2Kb restricted peptide #2.
 XX
 XX Ovalbumin-derived class I H-2Kb restricted peptide; vaccine;
 KW immunostimulant; immunogenic; interferon-gamma-producing T cell; IFN;
 KW popliteal lymph node; spleen; immune response; systemic response.
 XX
 OS Synthetic.
 XX
 PN WO200178767-A2.
 XX
 PD 25-OCT-2001.
 XX
 PR 17-APR-2001; 2001WO-EP04313.
 XX
 PR 14-APR-2000; 2000AT-0000657.
 XX
 PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
 XX
 PI Matzner F, Zauner W, Schmidt W, Buschle M;
 XX
 DR WPI; 2002-025970/03.
 XX
 PT Pharmaceutical preparation for use as a potent vaccine for inducing an
 PT improved immune response in a mammal, comprises a modified peptide -
 PT
 XX
 PS Example 1; Page 9; 18pp; English.
 XX
 CC The invention relates to a pharmaceutical preparation comprising a
 CC modified peptide, which induces an improved immune response in a mammal
 CC compared to the wild type peptide. The neutral peptide (SIINFEKL) (one
 CC negatively charged (Glu), one positively charged (Lys) amino acid) was
 CC rendered negative by adding (at the N-terminus) Glu Glu or Glu Asp Glu
 CC Asp, respectively. Results showed that the addition of 4 negatively-
 CC charged amino acids (EDBD) at the N-terminus of peptide SIINFEKL makes
 CC this peptide (in combination with poly-L-arginine) able to induce a high
 CC amount of specific interferon (IFN)-gamma-producing T cells in the
 CC draining (popliteal) lymph node (local response) and in the spleen
 CC (systemic response). Thus, the addition of hydrophobic amino acids as
 CC well as the addition of negatively charged amino acids transforms the
 CC peptide SIINFEKL to a good inducer of specific T cells. The modified
 CC peptides of the pharmaceutical composition induce a stronger immune
 CC response in a mammal compared to wild type antigens. The present
 CC sequence represents modified ovalbumin-derived class I H-2Kb restricted
 CC peptide #2 as described in the method of the invention.
 SQ Sequence 12 AA;

Query Match 100.0%; Score 38; DB 23; Length 12;
 Best Local Similarity 100.0%; Pred. No. 0.18;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 |||||
 DB 5 SIINFEKL 12

RESULT 82

AAU09826
 ID AAU09826 standard; peptide; 12 AA.

XX
 AC AAU09826;

DT 14-FEB-2002 (first entry)

XX Modified ovalbumin-derived class I H-2Kb restricted peptide #6.

XX Ovalbumin-derived class I H-2Kb restricted peptide; vaccine;

KW immunostimulant; immunogenic; interferon-gamma-producing T cell; IFN;

KW popliteal lymph node; spleen; immune response; systemic response.

OS Synthetic.

PN WO200178767-A2.

XX 25-OCT-2001.

XX 17-APR-2001; 2001WO-EP04313.

XX 14-APR-2000; 2000AT-0000657.

XX (CIST-) CISTEM BIOTECHNOLOGIES GMBH.

PI Matner F, Zauner W, Schmidt W, Buschle M;

DR WPI; 2002-025970/03.

PT Pharmaceutical preparation for use as a potent vaccine for inducing an
 improved immune response in a mammal, comprises a modified peptide -
 PS Example 2; Page 9; 18pp; English.

XX The invention relates to a pharmaceutical preparation comprising a
 CC modified peptide, which induces an improved immune response in a mammal
 CC compared to the wild type peptide. The neutral peptide (SIINFEKL) (one
 CC negatively charged (Glu), one positively charged (Lys) amino acid) was
 CC rendered negatively charged (at the N-terminus) Glu Glu or Glu Asp Glu
 CC Asp, respectively. Results showed that the addition of 4 negatively-
 CC charged amino acids (EBED) at the N-terminus of peptide SIINFEKL makes
 CC this peptide (in combination with poly-L-arginine) able to induce a high
 CC amount of specific interferon (IFN)-gamma-producing T cells in the
 CC draining (popliteal) lymph node (local response) and in the spleen
 CC (systemic response). Thus, the addition of hydrophobic amino acids as
 CC well as the addition of negatively charged amino acids transforms the
 CC peptide SIINFEKL to a good inducer of specific T cells. The modified
 CC peptides of the pharmaceutical composition induce a stronger immune
 CC response in a mammal compared to wild type antigens. The present
 CC sequence represents modified ovalbumin-derived class I H-2Kb restricted
 CC peptide #6 as described in the method of the invention.

XX Sequence 12 AA;

Query Match 100.0%; Score 38; DB 23; Length 12;
 Best Local Similarity 100.0%; Pred. No. 0.18;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 |||||
 DB 5 SIINFEKL 12

RESULT 83
 AAU09827
 ID AAU09827 standard; peptide; 12 AA.

XX
 AC AAU09827;

DT 14-FEB-2002 (first entry)

XX Modified ovalbumin-derived class I H-2Kb restricted peptide #7.

XX Ovalbumin-derived class I H-2Kb restricted peptide; vaccine;

KW immunostimulant; immunogenic; interferon-gamma-producing T cell; IFN;

KW popliteal lymph node; spleen; immune response; systemic response.

OS Synthetic.

PN WO200178767-A2.

XX 25-OCT-2001.

XX 17-APR-2001; 2001WO-EP04313.

XX 14-APR-2000; 2000AT-0000657.

XX (CIST-) CISTEM BIOTECHNOLOGIES GMBH.

PI Matner F, Zauner W, Schmidt W, Buschle M;

DR WPI; 2002-025970/03.

PT Pharmaceutical preparation for use as a potent vaccine for inducing an
 improved immune response in a mammal, comprises a modified peptide -
 PS Example 3; Page 10; 18pp; English.

XX The invention relates to a pharmaceutical preparation comprising a
 CC modified peptide, which induces an improved immune response in a mammal
 CC compared to the wild-type peptide. The neutral peptide (SIINFEKL) (one
 CC negatively charged (Glu), one positively charged (Lys) amino acid) was
 CC rendered negative by adding (at the N-terminus) Glu Glu or Glu Asp Glu
 CC Asp, respectively. Results showed that the addition of 4 negatively-
 CC charged amino acids (EBED) at the N-terminus of peptide SIINFEKL makes
 CC this peptide (in combination with poly-L-arginine) able to induce a high
 CC amount of specific interferon (IFN)-gamma-producing T cells in the
 CC draining (popliteal) lymph node (local response) and in the spleen
 CC (systemic response). Thus, the addition of hydrophobic amino acids as
 CC well as the addition of negatively charged amino acids transforms the
 CC peptide SIINFEKL to a good inducer of specific T cells. The modified
 CC peptides of the pharmaceutical composition induce a stronger immune
 CC response in a mammal compared to wild type antigens. The present
 CC sequence represents modified ovalbumin-derived class I H-2Kb restricted
 CC peptide #7 as described in the method of the invention.

XX Sequence 12 AA;

Query Match 100.0%; Score 38; DB 23; Length 12;
 Best Local Similarity 100.0%; Pred. No. 0.18;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 |||||
 DB 1 SIINFEKL 8

RESULT 84
 ABB76049
 ID ABB76049 standard; Peptide; 14 AA.

XX
 AC ABB76049;

DT 12-JUL-2002 (first entry)

DE Peptide insert in CACTES-Cys-Ova.
 XX Bordetella pertussis; adenylate cyclase; CyaA; adenylcyclase;
 KM vector; drug delivery; antigen delivery; cell targeting; CD11b;
 XX CACTES-Cys-Ova.
 OS Synthetic.
 XX EPI188446-A1.
 XX 20-MAR-2002.
 PD 15-SEP-2000; 2000EP-0402562.
 PF 15-SEP-2000; 2000EP-0402562.
 XX 15-SEP-2000; 2000EP-0402562.
 PR (INSP) INST PASTEUR.
 XX (CNRS) CENT NAT RECH SCT.
 PA Leduc C, Guernonprez P, Ladant D, Guiso N, Khelaf N;
 XX WPI; 2002-354020/39.
 DR Use of Bordetella adenylcyclase to make proteinaceous vector, useful
 XX for drug or antigen delivery, selectively targets cells that express
 PT CD11b -
 XX Example A; Page 10; 34pp; English.
 XX The present sequence is a peptide that was introduced into the
 CC catalytic domain of a detoxified form of the adenylate cyclase
 CC (CyaA, or adenylcyclase) of Bordetella pertussis. A recombinant
 CC detoxified CyaA toxin, CACTES-Cys-Ova, harbouring a unique cysteine
 CC that was derived from the peptide insert, was produced. The
 CC protein was labeled on its unique cysteine, and used to detect
 CC CyaA binding to neutrophils. Experiments showed that CyaA binding
 CC to the surface of 3 myeloid cell lines of mouse or human origin, as
 CC well as to human neutrophils, was mainly mediated through the
 CC CD11b/CD18 integrin. The invention relates to the novel use of
 CC Bordetella CyaA as a proteinaceous vector for targeting a
 CC molecule of interest to the surface CD11b-expressing cells,
 CC especially dendritic cells and neutrophils. The molecule of
 CC interest is translocated in the cytosol to prime a cytotoxic T
 CC lymphocyte response. In a preferred embodiment, a peptide is
 CC inserted into the catalytic domain of CyaA at a permissive site.
 CC The peptide may be an intracellular bacterial cell, tumour, viral,
 CC fungal or parasite cell antigen (all claimed). Alternatively, a
 CC drug, especially an antiinflammatory, is chemically coupled to
 CC CyaA for drug delivery.
 XX SQ Sequence 14 AA;
 QY Query Match 100.0%; Score 38; DB 23; Length 14;
 XX Best Local Similarity 100.0%; Pred. No. 0.22;
 DB Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEKL 8
 XX |||||
 DB 5 SIINFEKL 12
 RESULT 85
 AAU09823
 ID AAU09823 standard; peptide; 14 AA.
 XX
 AC AAU09823;
 XX
 DT 14-FEB-2002 (first entry)
 XX
 DE Modified ovalbumin-derived class I H-2Kb restricted peptide #3.
 XX
 KW Ovalbumin-derived class I H-2Kb restricted peptide; vaccine;
 KM immunostimulant; immunogenic; interferon-gamma-producing T cell; IFN;
 KW

KM popliteal lymph node; spleen; immune response; systemic response.
 XX Synthetic.
 OS
 XX WO200178767-A2.
 XX
 PD 25-OCT-2001.
 PF 17-APR-2001; 2001WO-EP04313.
 XX
 PR 14-APR-2000; 2000AT-0000657.
 XX
 PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
 XX
 PI Matner F, Zauner W, Schmidt W, Buschle M;
 XX WPI; 2002-025970/03.
 DR
 XX
 PT Pharmaceutical preparation for use as a potent vaccine for inducing an
 PT improved immune response in a mammal, comprises a modified peptide -
 XX
 XX Example 1; Page 9; 18pp; English.
 PS
 XX The invention relates to a pharmaceutical preparation comprising a
 CC modified peptide, which induces an improved immune response in a mammal
 CC compared to the wild type peptide. The neutral peptide (SIINFEKL) (one
 CC negatively charged (Glu), one positively charged (Lys) amino acid) was
 CC rendered negative by adding (at the N-terminus) Glu Glu or Glu Asp Glu
 CC Asp, respectively. Results showed that the addition of 4 negatively-
 CC charged amino acids (EDED) at the N-terminus of peptide SIINFEKL makes
 CC this peptide (in combination with poly-L-arginine) able to induce a high
 CC amount of specific interferon (IFN)-gamma-producing T cells in the
 CC draining (popliteal) lymph node (local response) and in the spleen
 CC (systemic response). Thus, the addition of hydrophobic amino acids as
 CC well as the addition of negatively charged amino acids transforms the
 CC peptide SIINFEKL to a good inducer of specific T cells. The modified
 CC peptides of the pharmaceutical composition induce a stronger immune
 CC response in a mammal compared to wild type antigens. The present
 CC sequence represents modified ovalbumin-derived class I H-2Kb restricted
 CC peptide #3 as described in the method of the invention.
 XX SQ Sequence 14 AA;
 QY Query Match 100.0%; Score 38; DB 23; Length 14;
 XX Best Local Similarity 100.0%; Pred. No. 0.22;
 DB Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEKL 8
 XX |||||
 DB 7 SIINFEKL 14
 RESULT 86
 AAU09828
 ID AAU09828 standard; peptide; 14 AA.
 XX
 AC AAU09828;
 XX
 DT 14-FEB-2002 (first entry)
 XX
 DE Modified ovalbumin-derived class I H-2Kb restricted peptide #8.
 XX
 KW Ovalbumin-derived class I H-2Kb restricted peptide; vaccine;
 KM immunostimulant; immunogenic; interferon-gamma-producing T cell; IFN;
 KW popliteal lymph node; spleen; immune response; systemic response.
 XX
 OS Synthetic.
 XX WO200178767-A2.
 XX
 PD 25-OCT-2001.
 PF 17-APR-2001; 2001WO-EP04313.
 XX

XX 14-APR-2000; 2000AT-0000657.
PR
XX
PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH.
XX
PI Mattern F, Zauner W, Schmidt W, Buschle M,
XX
XX WPI, 2002-025970/03.
DR
XX
PT Pharmaceutical preparation for use as a potent vaccine for inducing an
PT improved immune response in a mammal, comprises a modified peptide -
XX
XX
PS Example 4; Page 11; 18pp; English.

The invention relates to a pharmaceutical preparation comprising a modified peptide, which induces an improved immune response in a mammal compared to the wild type peptide. The neutral peptide (SIINFEKL) (one negatively charged (Glu), one positively charged (Lys) amino acid) was rendered negative by adding (at the N-terminus) Glu Glu or Glu Asp Glu Asp, respectively. Results showed that the addition of 4 negatively-charged amino acids (EEDD) at the N-terminus of peptide SIINFEKL makes this peptide (in combination with poly-L-arginine) able to induce a high amount of specific interferon (IFN)-gamma-producing T cells in the draining (popliteal) lymph node (local response) and in the spleen (systemic response). Thus, the addition of hydrophobic amino acids as well as the addition of negatively charged amino acids transforms the peptide SIINFEKL to a good inducer of specific T cells. The modified peptides of the pharmaceutical composition induce a stronger immune response in a mammal compared to wild type antigens. The present sequence represents modified ovalbumin-derived class I H-2Kb restricted peptide #8 as described in the method of the invention.

SQ Sequence 14 AA;

Query Match	100.0%	Score 38;	DB 23;	Length 14;
Best Local Similarity	100.0%;	Pred. No. 0.22;		
Matches	8;	Conservative	0;	Indels 0;
		Mismatches	0;	Gaps 0;

QY	1	SIINFEXL	8
Db	7	SIINFEXL	14

RESULT 87

AAU09824
ID AAU09824 standard; peptide; 15 AA.

AC AAU09824;

DT 14-FEB-2002 (first entry)

DE Modified ovalbumin-derived class I H-2Kb restricted peptide #4.

KW Ovalbumin-derived class I H-2Kb restricted peptide; vaccine; 11

popliteal lymph node; immune response; systemic response.

Synthetic.

PN WO200178767-A2.

PD 25-OCT-2001.

PF 17-APR-2001; 2001WO-EP04313.

PR 14-APR-2000; 2000AT-0000657.

PA (CIST-) CISTEM BIOTECHNOLOGIES GMBH

PI Mattner F, Zauner W, Schmidt W, Buschle M;

DR WPI; 2002-025970/03.

PT Pharmaceutical preparation for use as a potent vaccine for inducing an
PT improved immune response in a mammal, comprises a modified peptide -
XX
PS Example 1, Page 9, 18pp, English.

PS Example 1; Page 9; 18pp; English.

CC The invention relates to a pharmaceutical preparation comprising a
CC modified peptide, which induces an improved immune response in a mammal
CC compared to the wild type peptide. The neutral peptide (SIINFEKL) (one
CC negatively charged (Glu), one positively charged (Lys) amino acid) was
CC rendered negative by adding (at the N-terminus) Glu Glu or Glu Asp Glu
CC Asp, respectively. Results showed that the addition of 4 negatively-
CC charged amino acids (DDED) at the N-terminus of peptide SIINFEKL makes
CC this peptide (in combination with poly-DL-arginine) able to induce a high
CC amount of specific interferon (IFN)-gamma-producing T cells in the
CC draining (popliteal) lymph node (local response) and in the spleen
CC (systemic response). Thus, the addition of hydrophobic amino acids as
CC well as the addition of negatively charged amino acids transforms the
CC peptide SIINFEKL to a good inducer of specific T cells. The modified
CC peptides of the pharmaceutical composition induce a stronger immune
CC response in a mammal compared to wild type antigens. The present
CC sequence represents modified ovalbumin-derived class I H-2Kb restricted
CC peptide #4 as described in the method of the invention.

SQ Sequence 15 AA;

Query Match	100.0%;	Score 38;	DB 23;	Length 15;
Best Local Similarity	100.0%;	Pred. No. 0.23;	0;	Indels 0
Matches	8;	Conservative	0;	Mismatches 0;

Qy	1	SIINFEXL	8
Db	8	SIINFEXL	15

RESULT 88

ABP57403
ID ABP57403 standard; peptide; 16 AA

AC ABP57403;

DT 23-APR-2003 (first entry)

DE Synthetic 16mer peptide

KW Escherichia coli heat labile enterotoxin; Etx; cholera toxin; Ctx;

KW *Vibrio cholerae*; mutant; GM-1 ganglioside receptor; carrier; toxin;
 KW virucide; cytostatic; vaccine; viral infection; cancer; EtXB; CtxB.

OS Synthetic.

PN W02003000899-A1.

PD 03-JAN-2003

PF 20-JUN-2002; 2002WO-GB02829.

PR 22-JUN-2001; 2001GB-0015382.

PA (UYBR-) UNIV BRISTOL.

PI Hirst TR;

DR WPI; 2003-175291/17.

PT Use of a mutant form of B subunit of *Escherichia coli* heat labile

PT enterotoxin or B subunit of cholera toxin for delivering an agent to a target cell for treating viral infection or cancer -

PS Example 5; Page 45; 84pp; English.

2000 The present invention describes a mutant form of B subunit of Escherichia
2001 CC coli heat labile enterotoxin (EtxB) or B subunit of cholera toxin (CtxB)
2002 CC from *Vibrio cholerae* which is useful for delivering an agent to a target

cell, and has GM-1 ganglioside receptor binding activity but has reduced immunogenic and immunomodulatory activity relative to the wild-type form of Ectx or CtxB. Also described: (1) treating a disease or condition in a subject; (2) delivering the agent using the mutant to a target cell; (3) a composition; and (4) a kit for delivering the agent to a target cell. Mutant Ectx and CtxB have virucide and cytostatic activities and can be used in vaccines. The mutant can be used for the preparation of a medicament for delivering an exogenous peptide, which is the agent, into the major histocompatibility complex (MHC) Class I antigen processing and presenting pathways to elicit a cytotoxic T lymphocyte (CTL) response, or for separate, simultaneous or combined use for treating viral infection or cancer. The mutant form of Ectx or CtxB enters mammalian cells without inducing a potent anti-B-subunit response and immunomodulatory response. It may be linked with an agent to upregulate the presentation of the antigen or antigenic determinant. CC The present sequence represents a peptide which is used in an example from the present invention.

CC XX Sequence 16 AA;

Query Match 100.0%; Score 38; DB 24; Length 16;
Best Local Similarity 100.0%; Pred. No. 0.25; Mismatches 0; Indels 0; Gaps 0;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEXL 8
Db 9 SIINFEXL 16

RESULT 89

AAW19957
ID AAW19957 standard; Peptide; 19 AA.

AC AAW19957;

XX 10-NOV-1997 (first entry)

DT BiP-binding domain-OVA hybrid peptide.

DE BiP-binding domain-OVA hybrid peptide.

XX Vaccine; immunotherapy; heat shock protein; BiP; OVA; cancer;

KW infectious disease.

XX Synthetic.

OS Key

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

XX Hybrid peptides OVA-BiP (AAW19956) and BiP-OVA (AAW19957) comprise chicken OVA-peptide (see AAW19955) joined via a peptide linker to heat shock protein (HSP) BiP binding domain (see also AAW19951). CC The hybrid peptide can be combined in vitro with a HSP, such as hsp70, to form a complex that, when administered to a subject, induces an immune response.

CC XX Sequence 19 AA;

Query Match 100.0%; Score 38; DB 18; Length 19;
Best Local Similarity 100.0%; Pred. No. 0.3; Mismatches 0; Indels 0; Gaps 0;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEXL 8
Db 12 SIINFEXL 19

RESULT 90

AAW19956
ID AAW19956 standard; Peptide; 19 AA.

AC AAW19956;

XX 10-NOV-1997 (first entry)

DT OVA-BiP-binding domain hybrid peptide.

DE Vaccine; immunotherapy; heat shock protein; BiP; OVA; cancer;

KW infectious disease.

XX Synthetic.

OS Key

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

FT Peptide

Sequence 19 AA;

Example 1; Page 18; 58pp; English.

Query Match 100.0%; Score 38; DB 18; Length 19;
 Best Local Similarity 100.0%; Pred. No. 0.3;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEXL 8
 |||||
 DB 1 SIINFEXL 8

RESULT 91

AAE13446
 ID AAE13446 standard; peptide, 19 AA.

AC AAE13446;

DT 12-FEB-2002 (first entry)

XX Chicken MHC class I peptide antigen #1.

XX Antigenic complex; epitope; heat shock protein; HSP; tether; javelin;

KW major histocompatibility complex; MHC; therapy; immune response;

XX malignancy; chicken.

OS Gallus gallus.

XX Key Location/Qualifiers

FT Region 1..8 /note= "MHC class I epitope"

FT Region 9..11 /note= "Linker"

FT Region 12..19 /note= "Javelin sequence"

PN WO200179259-A1.

PD 25-OCT-2001.

PF 17-APR-2001; 2001WO-US12567.

PR 17-APR-2000; 2000US-197462P.

XX (ROTH/) ROTHMAN J E.

PA (MAYH/) MAYHEW M.

PA (HOEW/) HOE M.

PI Rothman JE, Mayhew M, Hoe M;

XX WPI; 2002-017594/02.

XX A new antigenic complex comprising epitopes non-covalently joined to a

PT heat shock protein by a molecular tether designated a javelin are

PT useful to treat or prevent infectious disease or malignancy -

XX Disclosure; Page 13; 47pp; English.

PS The present invention relates to an antigenic complex, comprising a

CC number of epitopes non-covalently joined to a heat shock protein (HSP) by

CC a tethering molecule referred to as javelin which has affinity for the

CC HSP under physiological conditions, where the epitopes are covalently

CC joined to the tethering molecule and one epitope is major

CC histocompatibility complex class I (MHC) and the other MHC class II. The

CC antigenic complex is used to induce immune responses directed towards the

CC treatment or prevention of infectious diseases and malignancies. The

CC present sequence is chicken MHC class I peptide antigen.

CC

XX

SQ Sequence 19 AA;

Query Match 100.0%; Score 38; DB 23; Length 19;

Best Local Similarity 100.0%; Pred. No. 0.3;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEXL 8
 |||||

DB 1 SIINFEXL 8

RESULT 92

AAE13447
 ID AAE13447 standard; peptide, 19 AA.

AC AAE13447;

DT 12-FEB-2002 (first entry)

XX Chicken MHC class I peptide antigen #2.

XX Antigenic complex; epitope; heat shock protein; HSP; tether; javelin;

KW major histocompatibility complex; MHC; therapy; immune response;

XX malignancy; chicken.

OS Gallus gallus.

XX Key Location/Qualifiers

FT Region 1..8 /note= "Javelin sequence"

FT Region 9..11 /note= "Linker"

FT Region 12..19 /note= "MHC class I epitope"

PN WO200179259-A1.

PD 25-OCT-2001.

PF 17-APR-2001; 2001WO-US12567.

PR 17-APR-2000; 2000US-197462P.

XX (ROTH/) ROTHMAN J E.

PA (MAYH/) MAYHEW M.

PA (HOEW/) HOE M.

PI Rothman JE, Mayhew M, Hoe M;

XX WPI; 2002-017594/02.

XX A new antigenic complex comprising epitopes non-covalently joined to a

PT heat shock protein by a molecular tether designated a javelin are

PT useful to treat or prevent infectious disease or malignancy -

XX Disclosure; Page 13; 47pp; English.

PS The present invention relates to an antigenic complex, comprising a

CC number of epitopes non-covalently joined to a heat shock protein (HSP) by

CC a tethering molecule referred to as javelin which has affinity for the

CC HSP under physiological conditions, where the epitopes are covalently

CC joined to the tethering molecule and one epitope is major

CC histocompatibility complex class I (MHC) and the other MHC class II. The

CC antigenic complex is used to induce immune responses directed towards the

CC treatment or prevention of infectious diseases and malignancies. The

CC present sequence is chicken MHC class I peptide antigen.

CC

XX

SQ Sequence 19 AA;

Query Match 100.0%; Score 38; DB 23; Length 19;

Best Local Similarity 100.0%; Pred. No. 0.3;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEXL 8
 |||||

DB 12 SIINFEXL 19

RESULT 93

ABP57404

ID ABP57404 standard; peptide, 19 AA.


```

XX AC ABP57404;
XX XX
XX DT 23-APR-2003 (first entry)
XX DE Synthetic 19mer peptide.
XX DE
XX KW Escherichia coli heat labile enterotoxin; Etx; cholera toxin; Ctx;
XX KW Vibrio cholerae; mutant; GM-1 ganglioside receptor; carrier; toxin;
XX KW virucide; cytostatic; vaccine; viral infection; cancer; EtxB; CtxB.
XX OS
XX OS Synthetic.
XX PN WO2003000899-A1.
XX XX
XX XX 03-JAN-2003.
XX PD
XX PF 20-JUN-2002; 2002WO-GB02829.
XX PR
XX PR 22-JUN-2001; 2001GB-0015382.
XX PA
XX PA (UYBR-) UNITV BRISTOL.
XX PI
XX PI Hirtel TR;
XX DR
XX DR WPI; 2003-175291/17.
XX PT
XX PT Use of a mutant form of B subunit of Escherichia coli heat labile
XX PT enterotoxin or B subunit of cholera toxin for delivering an agent to a
XX PT target cell for treating viral infection or cancer
XX PS
XX PS Example 5; Page 45; 84pp; English.
XX CC
XX CC The present invention describes a mutant form of B subunit of Escherichia
XX CC coli heat labile enterotoxin (EtxB) or B subunit of cholera toxin (CtxB)
XX CC from Vibrio cholerae which is useful for delivering an agent to a target
XX CC cell, and has GM-1 ganglioside receptor binding activity but has reduced
XX CC immunogenic and immunomodulatory activity relative to the wild-type form
XX CC of EtxB or CtxB. Also described: (1) treating a disease or condition in
XX CC a subject; (2) delivering the agent using the mutant to a target cell;
XX CC (3) a composition; and (4) a kit for delivering the agent to a target
XX CC cell. Mutant EtxB and CtxB have virucide and cytostatic activities and
XX CC can be used in vaccines. The mutant can be used for the preparation of
XX CC a medicament for delivering an exogenous peptide, which is the agent,
XX CC into the major histocompatibility complex (MHC) Class I antigen
XX CC processing and presenting pathways to elicit a cytotoxic T lymphocyte
XX CC (CTL) response, or for separate, simultaneous or combined use for
XX CC treating viral infection or cancer. The mutant form of EtxB or CtxB
XX CC enters mammalian cells without inducing a potent anti-B-subunit response
XX CC and immunomodulatory response. It may be linked with an agent to
XX CC upregulate the presentation of the antigen or antigenic determinant.
XX CC The present sequence represents a peptide which is used in an example
XX CC from the present invention.
XX SQ
XX SQ Sequence 19 AA;

```

```

Query Match 100.0%; Score 38; DB 24; Length 19;
Best Local Similarity 100.0%; Pred. No. 0.3;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 1 SIINFEKL 8
    |||||
Db 12 SIINFEKL 19

```

```

RESULT 94
AAR32294
ID AAR32294 standard; protein; 24 AA.
XX
XX AAR32294;
AC
XX
XX 25-MAR-2003 (updated)
DT 31-MAY-1993 (first entry)

```

```

XX DE Sequence of synthetic peptide ova 253-276 which corresp. to AAs 253-
XX DE 276 of ovalbumin.
XX DE
XX KW Cytotoxic T lymphocyte response; epitope; antigen.
XX KW
XX OS Synthetic.
XX OS
XX PN WO9301831-A1.
XX PD
XX PD 04-FEB-1993.
XX PF
XX PF 24-JUL-1992; 92WO-US06193.
XX PR
XX PR 25-JUL-1991; 91US-0735069.
XX PA
XX PA (IDEC-) IDEC PHARM CORP.
XX PI
XX PI Rastetter WH, Raychaudhuri S;
XX DR
XX DR WPI; 1993-058526/07.
XX PT
XX PT New compsn. comprising an antigen and a formulation - to induce a
XX PT cytotoxic T-lymphocyte response, useful for treating malaria,
XX PT HIV, influenza, hepatitis, herpes, cancer, etc.
XX PS
XX PS Disclosure; Page 19; 56pp; English.
XX CC
XX CC Carbone and Bevan demonstrated that cytotoxic T-lymphocyte (CTL)
XX CC induced in C57BL/6 mice by EGT-ova transfectant, and by
XX CC cytoplasmically ova-loaded splenocytes recognise E14 cells coated
XX CC with the peptide ova 258-276. To determine whether soluble ovalbumin
XX CC in AF induces similar CTL responses, spleen cells were prepared from
XX CC immunised mice and stimulated in vitro with EGT-ova. The effectors
XX CC were tested against E14 cells coated with the peptide ova 253-276
XX CC or with a control peptide derived from myelin basic protein (MBP 84-
XX CC 102). The results demonstrate that ova-AF primed CTL with a similar
XX CC specificity to those primed by transfectants, or by cytoplasmically
XX CC loaded ova.
XX CC (Updated on 25-MAR-2003 to correct PN field.)
XX SQ
XX SQ Sequence 24 AA;

```

```

Query Match 100.0%; Score 38; DB 14; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.38;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 1 SIINFEKL 8
    |||||
Db 5 SIINFEKL 12

```

```

RESULT 95
AAR41450
ID AAR41450 standard; Protein; 24 AA.
XX
XX AAR41450;
AC
XX
XX 25-MAR-2003 (updated)
DT 23-FEB-1994 (first entry)

```

```

DE Antigenic peptide bound by MHC class one molecules.

```

```

KW HLA; Human Leucocyte Antigen; MHC; Class one molecules; cancer;
KW autoimmunity; transplant rejection; T-cell activation.
XX
XX OS Synthetic.
XX OS
XX PN WO9317095-A1.
XX PD
XX PD 02-SEP-1993.
XX PF
XX PF 18-FEB-1993; 93WO-US01557.

```

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XX 19-FEB-1992; 92US-0841662.
XX (SCRI ) SCRIPPS RES INST.
XX Jackson M, Langlade-demoyen P, Peterson PA;
XX WPI; 1993-288401/36.
XX Prodn. and use of human class I MHC molecules for activation of
XX CD8 cells - for therapy of e.g. cancer, viral, retroviral and
XX auto-immune diseases
XX
XX PS Disclosure; Page 77; 182pp; English.
XX
XX CC Human class I MHC genes are inserted into a cell and placed under
XX the control of an inducible promoter. This provides a means of
XX producing, loading and using Class I MHC molecules to specifically
XX activate CD8 cells in vitro. Activated cells can be used to
XX specifically kill target cells and also to treat cancer as well as
XX viral, retroviral, autoimmune and autoimmuno-type diseases. When
XX conjugated to a toxin, empty human MHC molecules expressed by the
XX cells can be used to inhibit transplant rejection. A number of
XX antigenic peptides (AAR41450-R41463) are synthesised to be bound by
XX the MHC molecules and this binding can then activate the CD8 cells.
XX (Updated on 25-MAR-2003 to correct PN field.)
XX
XX SQ Sequence 24 AA;

```

```

Query Match 100.0%; Score 38; DB 14; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.38;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY 1 STINFEKL 8
   |||||
DB 5 STINFEKL 12

```

```

RESULT 96
AAW04645
ID AAW04645 standard; peptide; 24 AA.
XX
XX AAW04645;
XX
XX 01-AUG-1997 (first entry)
XX
XX Ovalbumin-derived activated CD8+ T cells epitope OVA24.
XX
XX Macrophage; artificial antigen presenting cell; APC; cancer;
XX tumours; neoplasia; viral infection; retroviral infection;
XX autoimmune.
XX
XX Synthetic.
XX
XX WO9637107-A1.
XX
XX 28-NOV-1996.
XX
XX 22-MAY-1996; 96WO-US07436.
XX
XX 23-MAY-1995; 95US-0447761.
XX
XX (SCRI ) SCRIPPS RES INST.
XX
XX DeBruin MLH, Jackson MR, Peterson PA;
XX WPI; 1997-020850/02.
XX
XX Prodn. of activated CD8+ T cells directed to specific antigen - can
XX specifically kill target cells useful to treat, e.g. cancer
XX
XX Example 1; Page 26; 84pp; English.
XX

```

```

CC The method for the production of activated CD8+ T cells specifically
CC directed towards a particular antigen involves affixing peptides
CC corresponding to the particular antigen to an artificial support;
CC contacting macrophages with the affixed peptides for a time sufficient
CC for the peptides to be engulfed, and at least a portion of the peptides
CC to be presented on the surface of the macrophage; and contacting
CC unprimed CD8+ T cells with the peptide presenting macrophages for a
CC time sufficient to activate the unprimed CD8+ T cells. The present
CC sequence represents a peptide designated OVA24 which corresponds to
CC ovalbumin, a Kb-restricted peptide antigen. This is not as efficient as
CC the optimal peptide. The method, macrophages and artificial antigen
CC presenting cell, having a peptide corresponding to the particular
CC antigen present on its surface and at least a portion of an artificial
CC support in its interior, can be used to treat conditions (e.g. cancer,
CC tumours, neoplasia, viral or retroviral infection or autoimmune or
CC autoimmuno-type conditions) in patients via the specific killing of
CC target cells.

```

```

SQ Sequence 24 AA;

```

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Query Match 100.0%; Score 38; DB 18; Length 24;
Best Local Similarity 100.0%; Pred. No. 0.38;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY 1 STINFEKL 8
   |||||
DB 5 STINFEKL 12

```

```

RESULT 97
AAG65170
ID AAG65170 standard; peptide; 24 AA.
XX
XX AAG65170;
XX
XX 15-OCT-2002 (first entry)
XX
XX
XX Ovalbumin based peptide.
XX
XX Cytotoxic T-lymphocyte response; CTL; antigen; viral infection;
XX bacterial infection; cancer; parasitic infection; immune response;
XX non-toxic.
XX
XX Unidentified.
XX
XX US6270769-B1.
XX
XX 07-AUG-2001.
XX
XX 24-MAY-1995; 95US-0449728.
XX
XX 24-JUL-1992; 92US-0919787.
XX
XX 25-JUL-1991; 91US-0735069.
XX
XX (IDEC-) IDEC PHARM CORP.
XX
XX Raychaudhuri S, Rastetter WH;
XX WPI; 2001-564234/63.
XX
XX Induction of cytotoxic T-lymphocyte responses -
XX
XX Disclosure; Column 17-18; 24pp; English.
XX
XX The present invention relates to a method of treating viral, parasitic
XX and bacterial infections and cancer in humans, by administering an
XX antigen which causes a cytotoxic T-lymphocyte response. Said antigen does
XX not contain an immunostimulatory element. The method can also be used in
XX domesticated animals. The present sequence is a peptide used as an
XX antigen in the exemplification of the invention.
XX
XX SQ Sequence 24 AA;

```

Query Match 100.0%; Score 38; DB 22; Length 24;
 Best Local Similarity 100.0%; Pred. No. 0.38;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 DB 5 SIINFEKL 12

RESULT 98
 AAB74439
 ID AAB74439 standard; peptide; 24 AA.
 AC AAB74439;
 XX
 XX
 DT 29-MAY-2001 (first entry)
 XX
 DE Myelin basic protein amino acids 84-102 peptide.
 XX
 KW Cytotoxic T-lymphocyte; CTL response; immunostimulation; infection;
 XX cancer.
 OS Unidentified.
 XX
 PN US6197311-B1.
 XX
 PD 06-MAR-2001.
 XX
 PF 17-FEB-1998; 98US-0024220.
 XX
 PR 07-JUN-1995; 95US-0476674.
 XX
 PR 25-JUL-1991; 91US-0735069.
 PR 24-JUL-1992; 92US-0919787.
 PR 07-DEC-1994; 94US-0351001.
 XX
 PA (IDEC-) IDEC PHARM CORP.
 XX
 PI Raychaudhuri S, Rastetter WH, Black A;
 XX
 DR MPI; 2001-256350/26.
 XX
 PT Treating papillomavirus-related tumor or malignancy, involves
 PT administering an antigen formulation substantially free of
 PT immunostimulatory peptides, and comprising human papillomavirus antigen
 PT and microfluidized adjuvant -
 XX
 PS Disclosure; Column 10-11; 22pp; English.
 XX
 CC The present invention describes a method of treating a
 CC papillomavirus-related tumor, involving administering an antigen
 CC formulation capable of inducing a cytotoxic T-lymphocyte (CTL) response
 CC specific to the papillomavirus antigen in the individual. This is useful
 CC in the treatment of cancer and infections, such as those due to HIV,
 CC bacteria, parasites, influenza, herpes virus and hepatitis viruses. The
 CC present sequence is a peptide used to demonstrate the method of the
 CC invention.
 CC
 SQ Sequence 24 AA;

Query Match 100.0%; Score 38; DB 22; Length 24;
 Best Local Similarity 100.0%; Pred. No. 0.38;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 DB 5 SIINFEKL 12

RESULT 99
 ABG31664
 ID ABG31664 standard; Peptide; 24 AA.
 AC ABG31664;

XX 05-NOV-2002 (first entry)
 DT
 XX
 DE Ovalbumin (OVA) peptide fragment.
 KW Ovalbumin; microfluidised antigen; detergent; micelle-forming agent;
 KW biodegradable oil; biocompatible oil; cytotoxic T lymphocyte; HIV;
 KW human immunodeficiency virus; herpes virus; malaria; influenza; cancer;
 KW hepatitis; respiratory syncytial virus; domesticated animal; OVA;
 KW agricultural animal.
 XX
 OS Unidentified.
 XX
 PN US2002039582-A1.
 XX
 PD 04-APR-2002.
 XX
 PF 20-DEC-2000; 2000US-0740003.
 XX
 PR 17-FEB-1998; 98US-0024220.
 PR 25-JUL-1991; 91US-0735069.
 PR 07-DEC-1994; 94US-0351001.
 PR 07-JUN-1995; 95US-0476674.
 PR 29-AUG-1997; 97US-0919787.
 XX
 PA (IDEC-) IDEC PHARM CORP.
 XX
 PI Raychaudhuri S, Rastetter WH, Black A;
 XX
 DR MPI; 2002-607062/65.
 XX
 PT Composition useful for inducing cytotoxic T lymphocyte response in
 PT domesticated animals and humans comprises antigen mixed with
 PT microfluidised antigen formulation which is substantially free of
 PT immunostimulating peptides -
 XX
 PS Disclosure; Page 6; 31pp; English.
 XX
 CC The invention relates to a composition comprising an antigen mixed with
 CC microfluidised antigen formulation comprising a stabilised detergent, a
 CC micelle-forming agent and a biodegradable/biocompatible oil. The
 CC composition is formulated as a stable oil-in-water emulsion substantially
 CC free of or lacking immunostimulating peptides, and is capable of inducing
 CC specific cytotoxic T lymphocyte response against antigens in vivo. The
 CC composition is useful for treating patients infected with human
 CC immunodeficiency virus (HIV) or herpes virus, and patients suffering from
 CC malaria, influenza, hepatitis, cancer or respiratory syncytial virus by
 CC administering a composition comprising HIV antigen, malaria-associated
 CC antigen, hepatitis-associated antigen, cancer-associated antigen, herpes
 CC antigen or respiratory syncytial antigen respectively, mixed with
 CC microfluidised antigen consisting essentially of two of stabilising
 CC detergent, micelle-forming agent and biodegradable and biocompatible oil,
 CC the antigen formulation being formulated as stable oil-in-water emulsion,
 CC and inducing cytotoxic T lymphocyte response in a patient e.g. human,
 CC domesticated animal or agricultural animal. This sequence represents an
 CC ovalbumin (OVA) peptide fragment used in the scope of the invention.
 CC
 SQ Sequence 24 AA;

Query Match 100.0%; Score 38; DB 23; Length 24;
 Best Local Similarity 100.0%; Pred. No. 0.38;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 DB 5 SIINFEKL 12

RESULT 100
 ABP57405
 ID ABP57405 standard; peptide; 26 AA.
 AC ABP57405;

```

XX 23-APR-2003 (first entry)
XX Synthetic 26mer peptide.
XX
XX Escherichia coli heat labile enterotoxin; Etx; cholera toxin; Ctx;
XX Vibrio cholerae; mutant; GM-1 ganglioside receptor; carrier; toxin;
XX virucide; cytostatic; vaccine; viral infection; cancer; EtxB; CtxB.
XX
XX Synthetic.
XX
XX WO2003000899-A1.
XX
XX 03-JAN-2003.
XX
XX 20-JUN-2002; 2002WO-GB02829.
XX
XX 22-JUN-2001; 2001GB-0015382.
XX
XX (UYBR-) UNIV BRISTOL.
XX
XX Hirst TR;
XX
XX WPI; 2003-175291/17.
XX
XX Use of a mutant form of B subunit of Escherichia coli heat labile
XX enterotoxin or B subunit of cholera toxin for delivering an agent to a
XX target cell for treating viral infection or cancer -
XX
XX Example 5; Page 45; 84pp; English.
XX
XX The present invention describes a mutant form of B subunit of Escherichia
XX coli heat labile enterotoxin (EtxB) or B subunit of cholera toxin (CtxB)
XX from Vibrio cholerae which is useful for delivering an agent to a target
XX cell, and has GM-1 ganglioside receptor binding activity but has reduced
XX immunogenic and immunomodulatory activity relative to the wild-type form
XX of EtxB or CtxB. Also described: (1) treating a disease or condition in
XX a subject; (2) delivering the agent using the mutant to a target cell;
XX (3) a composition; and (4) a kit for delivering the agent to a target
XX cell. Mutant EtxB and CtxB have virucide and cytostatic activities and
XX can be used in vaccines. The mutant can be used for the preparation of
XX a medicament for delivering an exogenous peptide, which is the agent,
XX into the major histocompatibility complex (MHC) Class I antigen
XX processing and presenting pathways to elicit a cytotoxic T lymphocyte
XX (CTL) response, or for separate, simultaneous or combined use for
XX treating viral infection or cancer. The mutant form of EtxB or CtxB
XX enters mammalian cells without inducing a potent anti-B-subunit response
XX and immunomodulatory response. It may be linked with an agent to
XX upregulate the presentation of the antigen or antigenic determinant.
XX The present sequence represents a peptide which is used in an example
XX from the present invention.
XX
XX SQ Sequence 26 AA;
XX
XX Query Match 100.0%; Score 38; DB 24; Length 26;
XX Best Local Similarity 100.0%; Pred. No. 0.41;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 SIINFEXT 8
XX |||||
XX DB 19 SIINFEXT 26
XX
XX RESULT 101
XX ABP57406 standard; peptide; 26 AA.
XX
XX AC ABP57406;
XX
XX DT 23-APR-2003 (first entry)
XX
XX DE Synthetic 26mer* peptide.
XX
XX

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```

KW Escherichia coli heat labile enterotoxin; Etx; cholera toxin; Ctx;
KW Vibrio cholerae; mutant; GM-1 ganglioside receptor; carrier; toxin;
KW virucide; cytostatic; vaccine; viral infection; cancer; EtxB; CtxB.
KW
KW Synthetic.
KW
KW WO2003000899-A1.
KW
KW 03-JAN-2003.
KW
KW 20-JUN-2002; 2002WO-GB02829.
KW
KW 22-JUN-2001; 2001GB-0015382.
KW
KW (UYBR-) UNIV BRISTOL.
KW
KW Hirst TR;
KW
KW WPI; 2003-175291/17.
KW
KW Use of a mutant form of B subunit of Escherichia coli heat labile
KW enterotoxin or B subunit of cholera toxin for delivering an agent to a
KW target cell for treating viral infection or cancer -
KW
KW Example 5; Page 45; 84pp; English.
KW
KW The present invention describes a mutant form of B subunit of Escherichia
KW coli heat labile enterotoxin (EtxB) or B subunit of cholera toxin (CtxB)
KW from Vibrio cholerae which is useful for delivering an agent to a target
KW cell, and has GM-1 ganglioside receptor binding activity but has reduced
KW immunogenic and immunomodulatory activity relative to the wild-type form
KW of EtxB or CtxB. Also described: (1) treating a disease or condition in
KW a subject; (2) delivering the agent using the mutant to a target cell;
KW (3) a composition; and (4) a kit for delivering the agent to a target
KW cell. Mutant EtxB and CtxB have virucide and cytostatic activities and
KW can be used in vaccines. The mutant can be used for the preparation of
KW a medicament for delivering an exogenous peptide, which is the agent,
KW into the major histocompatibility complex (MHC) Class I antigen
KW processing and presenting pathways to elicit a cytotoxic T lymphocyte
KW (CTL) response, or for separate, simultaneous or combined use for
KW treating viral infection or cancer. The mutant form of EtxB or CtxB
KW enters mammalian cells without inducing a potent anti-B-subunit response
KW and immunomodulatory response. It may be linked with an agent to
KW upregulate the presentation of the antigen or antigenic determinant.
KW The present sequence represents a peptide which is used in an example
KW from the present invention.
KW
KW SQ Sequence 26 AA;
KW
KW Query Match 100.0%; Score 38; DB 24; Length 26;
KW Best Local Similarity 100.0%; Pred. No. 0.41;
KW Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
KW
KW QY 1 SIINFEXT 8
KW |||||
KW DB 19 SIINFEXT 26
KW
KW RESULT 102
KW AAE13448
KW ID AAE13448 standard; peptide; 30 AA.
KW
KW AC AAE13448;
KW
KW DT 12-FEB-2002 (first entry)
KW
KW DE Chicken MHC class I peptide antigen #3.
KW
KW AC Antigenic complex; epitope; heat shock protein; HSP; tether; javelin;
KW major histocompatibility complex; MHC; therapy; immune response;
KW malignancy; chicken.
KW
KW OS Gallus gallus.

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XX Key Location/Qualifiers
FH Region 1..8
FT /note= "Javelin sequence"
FT Region 9..11
FT /note= "Linker"
FT Region 12..19
FT /note= "MHC class I epitope"
FT Region 20..22
FT /note= "Linker"
FT Region 23..30
FT /note= "Javelin sequence"
XX
XX WO200179259-A1.
XX
XX 25-OCT-2001.
XX
XX 17-APR-2001; 2001WO-US12567.
XX
XX 17-APR-2000; 2000US-197462P.
XX
XX (ROTH/) ROTHMAN J E.
XX (MAYH/) MAYHEW M.
XX (HOEM/) HOE M.
XX
XX Rothman JF, Mayhew M, Hoe M;
XX
XX WPI; 2002-017594/02.
XX
XX A new antigenic complex comprising epitopes non-covalently joined to a
XX heat shock protein by a molecular tether designated a javelin are
XX useful to treat or prevent infectious disease or malignancy -
XX
XX Disclosure; Page 13; 47pp; English.
XX
XX The present invention relates to an antigenic complex, comprising a
XX number of epitopes non-covalently joined to a heat shock protein (HSP) by
XX a tethering molecule referred to as javelin which has affinity for the
XX HSP under physiological conditions, where the epitopes are covalently
XX joined to the tethering molecule and one epitope is major
XX histocompatibility complex class I (MHC) and the other MHC class II. The
XX antigenic complex is used to induce immune responses directed towards the
XX treatment or prevention of infectious diseases and malignancies. The
XX present sequence is chicken MHC class I peptide antigen.
XX
XX Sequence 30 AA;
XX
XX Query Match 100.0%; Score 38; DB 23; Length 30;
XX Best Local Similarity 100.0%; Pred. No. 0.48;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 SIINFEKL 8
XX |||||
XX DB 12 SIINFEKL 19
XX
XX RESULT 103
XX ABP57407
XX ID ABP57407 standard; peptide; 31 AA.
XX
XX ABP57407;
XX
XX 23-APR-2003 (first entry)
XX
XX Synthetic 31mer peptide.
XX
XX Escherichia coli heat labile enterotoxin; Etx; cholera toxin; Ctx;
XX Vibrio cholerae; mutant; GM-1 ganglioside receptor; carrier; toxin;
XX virucide; cytostatic; vaccine; viral infection; cancer; CtxB; CtxB.
XX Synthetic.
XX OS
XX WO2003000899-A1.
XX PN

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XX PD 03-JAN-2003.
XX
XX 20-JUN-2002; 2002WO-GB02829.
XX
XX 22-JUN-2001; 2001GB-0015382.
XX
XX (UYBR-) UNIV BRISTOL.
XX
XX Hirst TR;
XX
XX WPI; 2003-175291/17.
XX
XX Use of a mutant form of B subunit of Escherichia coli heat labile
XX enterotoxin or B subunit of cholera toxin for delivering an agent to a
XX target cell for treating viral infection or cancer -
XX
XX Example 5; Page 45; 84pp; English.
XX
XX The present invention describes a mutant form of B subunit of Escherichia
XX coli heat labile enterotoxin (EtxB) or B subunit of cholera toxin (CtxB)
XX from Vibrio cholerae which is useful for delivering an agent to a target
XX cell, and has GM-1 ganglioside receptor binding activity but has reduced
XX immunogenic and immunomodulatory activity relative to the wild-type form
XX of EtxB or CtxB. Also described: (1) treating a disease or condition in
XX a subject; (2) delivering the agent using the mutant to a target cell;
XX (3) a composition; and (4) a kit for delivering the agent to a target
XX cell. Mutant EtxB and CtxB have virucide and cytostatic activities and
XX can be used in vaccines. The mutant can be used for the preparation of
XX a medicament for delivering an exogenous peptide, which is the agent,
XX into the major histocompatibility complex (MHC) Class I antigen
XX processing and presenting pathways to elicit a cytotoxic T lymphocyte
XX (CTL) response, or for separate, simultaneous or combined use for
XX treating viral infection or cancer. The mutant form of EtxB or CtxB
XX enters mammalian cells without inducing a potent anti-B-subunit response
XX and immunomodulatory response. It may be linked with an agent to
XX upregulate the presentation of the antigen or antigenic determinant.
XX The present sequence represents a peptide which is used in an example
XX from the present invention.
XX
XX Sequence 31 AA;
XX
XX Query Match 100.0%; Score 38; DB 24; Length 31;
XX Best Local Similarity 100.0%; Pred. No. 0.49;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 SIINFEKL 8
XX |||||
XX DB 19 SIINFEKL 26
XX
XX RESULT 104
XX AAM04646
XX ID AAM04646 standard; peptide; 35 AA.
XX
XX AAM04646;
XX
XX 01-AUG-1997 (first entry)
XX
XX Ovalbumin-derived activated CD8+ T cells epitope OVA35.
XX
XX Macrophage; artificial antigen presenting cell; APC; cancer;
XX tumours; neoplasia; viral infection; retroviral infection;
XX autoimmune.
XX Synthetic.
XX OS
XX WO9637107-A1.
XX 28-NOV-1996.
XX
XX 22-MAY-1996; 96WO-US07436.
XX
XX

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PR 23-MAY-1995; 95US-0447761.
XX
XX (SCRI ) SCRIPPS RES INST.
XX
XX DeBrujn MLH, Jackson MR, Peterson PA;
XX
XX WPI; 1997-020850/02.
XX
XX Prodn. of activated CD8+ T cells directed to specific antigen - can
XX specifically kill target cells useful to treat, e.g. cancer
XX
XX Example 1; Page 26; 84pp; English.
XX
XX The method for the production of activated CD8+ T cells specifically
XX directed towards a particular antigen involves affixing peptides
XX corresponding to the particular antigen to an artificial support;
XX contacting macrophages with the affixed peptides for a time sufficient
XX for the peptides to be engulfed, and at least a portion of the peptides
XX to be presented on the surface of the macrophage; and contacting
XX unprimed CD8+ T cells with the peptide presenting macrophages for a
XX time sufficient to activate the unprimed CD8+ T cells. The present
XX sequence represents a peptide designated OVA35 which corresponds to
XX ovalbumin, a Kb-restricted peptide antigen. This is not as efficient as
XX the optimal peptide. The method, macrophages and artificial antigen
XX presenting cell, having a peptide corresponding to the particular
XX antigen present on its surface and at least a portion of an artificial
XX support in its interior, can be used to treat conditions (e.g. cancer,
XX tumours, neoplasia, viral or retroviral infection or autoimmune or
XX autoimmune-type conditions) in patients via the specific killing of
XX target cells.
XX
XX Sequence 35 AA;
SQ
Query Match 100.0%; Score 38; DB 18; Length 35;
Best Local Similarity 100.0%; Pred. No. 0.56;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 STINPEKL 8
DB 16 STINPEKL 23
RESULT 105
AAO26741
ID AAO26741 standard; peptide; 36 AA.
XX
XX AAO26741;
XX
XX 27-MAR-2003 (first entry)
XX
XX Chicken ovalbumin class I and II carrying antigenic peptide.
XX
XX Cytostatic; universal polypeptidic carrier; Gb3 receptor; cytotoxic drug;
XX tumour cell; immunogenic; gene therapy; chicken.
XX
XX Gallus sp.
XX
XX EPI229045-A1.
XX
XX 07-AUG-2002.
XX
XX 01-FEB-2001; 2001EP-0400255.
XX
XX 01-FEB-2001; 2001EP-0400255.
XX
XX (CUR1-) INST CURIE.
XX (CNRS ) CENT NAT RECH SCI.
XX (INRM ) INSERM INST NAT SANTE & RECH MEDICALE.
XX (UYPA-) UNIV CURIE PARIS VI P & M.
XX
XX Johannes L, Tartour E, Goud B, Fridman WH;
XX
XX WPI; 2003-185783/19.

```

```

XX
XX Novel universal polypeptidic carrier for targeting a molecule directly
XX or indirectly to Gb3 receptor expressing cells; useful for directing
XX cytotoxic drugs to tumor cells expressing Gb3 receptor -
XX
XX Example 3; Page 8; 24pp; English.
XX
XX The invention relates to a universal polypeptidic carrier for targeting
XX directly or indirectly a molecule to Gb3 receptor expressing cells. The
XX universal polypeptidic carrier is useful for targeting directly or
XX indirectly, to Gb3 receptor expressing cells, using molecules such as
XX proteins, peptides, oligopeptides, glycoproteins, glycopeptides, nucleic
XX acids, polynucleotides, or its combination, an antigen to be targeted
XX to antigen presenting cells or a cytotoxic drug or pro-drug to be targeted
XX to tumor cells expressing Gb3 receptor. The universal polypeptidic
XX carrier is useful for delivering an expression vector containing a
XX sequence of interest into a Gb3 receptor expressing cells which involves
XX operably linking the expression vector to a lysine-rich peptide
XX covalently linked to the Cys moiety of the universal polypeptidic
XX carrier. The lysine rich peptide is preferably a 16-mer poly-lysine and
XX the sequence of interest is preferably a sequence encoding an immunogenic
XX peptide, or a sequence encoding a drug or a pro-drug becoming toxic for
XX the Gb3 receptor expressing cells, or a sequence encoding a therapeutic
XX active molecule. The universal polypeptidic carrier is useful for
XX targeting a molecule to a Gb3 receptor expressing cell to enable the
XX molecule to be internalised, processed and/or expressed in the cell
XX expressing a Gb3 receptor. The universal polypeptidic carrier is useful
XX as a carrier for introducing a nucleotide sequence in a target cell
XX either for gene therapy or for obtaining recombinant cells expressing
XX heterologous proteins. This sequence represents a chicken ovalbumin class
XX I and II carrying antigenic peptide relating to the universal carrier
XX protein of the invention.
XX
XX Sequence 36 AA;
SQ
Query Match 100.0%; Score 38; DB 24; Length 36;
Best Local Similarity 100.0%; Pred. No. 0.58;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 STINPEKL 8
DB 6 STINPEKL 13
RESULT 106
AAB84325
ID AAB84325 standard; peptide; 43 AA.
XX
XX AAB84325;
XX
XX 22-AUG-2001 (first entry)
XX
XX Amino acid sequence of a lems variant peptide.
XX
XX lemA; CD8+ epitope; T cell response.
XX
XX Synthetic.
XX
XX WO200140275-A2.
XX
XX 07-JUN-2001.
XX
XX 06-DEC-2000; 2000WO-US33027.
XX
XX 06-DEC-1999; 99US-0169227.
XX
XX (USSH ) US DEPT HEALTH & HUMAN SERVICES.
XX
XX Kurlander RJ, Chao E, Fields J;
XX
XX WPI; 2001-389952/41.
XX
XX New isolated variant of lemA, comprising a hydrophobic element
XX

```

PT joined to a CD8+ epitope, useful for inducing a directed CD8+ T cell
 PT response or as a treatment or prophylactic against diseases -
 XX
 XX
 PS Disclosure; Page 17; 65pp; English.

CC The specification describes a peptide variant of IemA, comprising a
 CC hydrophobic element joined to a CD8+ epitope. The peptides may be
 CC used therapeutically by administering the peptides to a patient having
 CC a need to induce a directed CD8+ T cell response. The peptide may also
 CC be used as a preventive measure to avoid a disease or condition, or to
 CC treat subjects already afflicted with a disease. The present sequence
 CC represents a peptide of the invention.
 XX

SQ Sequence 43 AA;

Query Match 100.0%; Score 38; DB 22; Length 43;
 Best Local Similarity 100.0%; Pred. No. 0.69;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEXL 8
 |||||
 DB 28 SIINFEXL 35

RESULT 107
 AAB84321
 ID AAB84321 standard; peptide; 47 AA.

AC AAB84321;

DT 22-AUG-2001 (first entry)

DE Amino acid sequence of a IemA variant peptide.

KM IemA; CD8+ epitope; T cell response.

OS Synthetic.

PN WO200140275-A2.

PD 07-JUN-2001.

PF 06-DEC-2000; 2000WO-US33027.

PR 06-DEC-1999; 99US-0169227.

PA (USSH) US DEPT HEALTH & HUMAN SERVICES.

PI Kurlander RJ, Chao E, Fields J;

DR WPI; 2001-389952/41.

PT New isolated variant of IemA, comprising a hydrophobic element
 PT joined to a CD8+ epitope, useful for inducing a directed CD8+ T cell
 PT response or as a treatment or prophylactic against diseases -
 XX

PS Disclosure; Page 19; 65pp; English.

CC The specification describes a peptide variant of IemA, comprising a
 CC hydrophobic element joined to a CD8+ epitope. The peptides may be
 CC used therapeutically by administering the peptides to a patient having
 CC a need to induce a directed CD8+ T cell response. The peptide may also
 CC be used as a preventive measure to avoid a disease or condition, or to
 CC treat subjects already afflicted with a disease. The present sequence
 CC represents a peptide of the invention.
 XX

SQ Sequence 47 AA;

Query Match 100.0%; Score 38; DB 22; Length 47;
 Best Local Similarity 100.0%; Pred. No. 0.76;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEXL 8

DB |||||
 34 SIINFEXL 41

RESULT 108

AAB84322
 ID AAB84322 standard; peptide; 48 AA.

AC AAB84322;

DT 22-AUG-2001 (first entry)

DE Amino acid sequence of a IemA variant peptide.

KM IemA; CD8+ epitope; T cell response.

OS Synthetic.

PN WO200140275-A2.

PD 07-JUN-2001.

PF 06-DEC-2000; 2000WO-US33027.

PR 06-DEC-1999; 99US-0169227.

PA (USSH) US DEPT HEALTH & HUMAN SERVICES.

PI Kurlander RJ, Chao E, Fields J;

DR WPI; 2001-389952/41.

PT New isolated variant of IemA, comprising a hydrophobic element
 PT joined to a CD8+ epitope, useful for inducing a directed CD8+ T cell
 PT response or as a treatment or prophylactic against diseases -
 XX

PS Disclosure; Page 19; 65pp; English.

CC The specification describes a peptide variant of IemA, comprising a
 CC hydrophobic element joined to a CD8+ epitope. The peptides may be
 CC used therapeutically by administering the peptides to a patient having
 CC a need to induce a directed CD8+ T cell response. The peptide may also
 CC be used as a preventive measure to avoid a disease or condition, or to
 CC treat subjects already afflicted with a disease. The present sequence
 CC represents a peptide of the invention.
 XX

SQ Sequence 48 AA;

Query Match 100.0%; Score 38; DB 22; Length 48;
 Best Local Similarity 100.0%; Pred. No. 0.78;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEXL 8
 |||||
 DB 1 SIINFEXL 8

RESULT 109

AAB48953
 ID AAB48953 standard; Protein; 49 AA.

AC AAB48953;

DT 27-MAR-2001 (first entry)

DE Tn5-DICE ovalbumin MHC class I epitope fusion protein.

KM Transposable element; MHC epitope; major histocompatibility complex;
 KM intracellular bacterial pathogen; IxP site; Cre recombinase;
 KM insertion end; In-frame fusion; detection; antigen;
 KM disseminated insertions of class-I epitopes; DICE-I; transposon Tn5;
 KM ovalbumin MHC class I epitope.
 XX

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OS Synthetic.
XX
XX WO200071158-A1.
XX
XX 30-NOV-2000.
XX
XX 26-MAY-2000; 2000WO-US14687.
XX
XX 26-MAY-1999; 99US-0136210.
XX
XX (UYOR-) UNIV OREGON HEALTH SCI.
XX
XX Heffron FL, Parker DC, Ellefson DD;
XX
XX MPI; 2001-031967/04.
XX
XX Transposable element for detecting an antigenic epitope of a pathogen,
XX comprising 5' and 3' recombining sites, nucleic acid sequences encoding
XX a selectable marker and major histocompatibility complex (MHC) epitope,
XX and an insertion end -
XX
XX Example 2; Fig 2; 63pp; English.
XX
XX The invention relates to a novel transposable element comprising DNA
XX encoding a selectable marker (e.g., antibiotic resistance) located
XX between a 5' recombining site and a 3' recombining site (e.g., loxp
XX sites); DNA encoding an MHC (major histocompatibility complex) epitope
XX either 5' of the 5' recombining site or 3' of the 3' recombining site;
XX and insertion ends comprising an inverted repeat sequence at the 5' and
XX 3' ends of the transposable element sufficient for integration of the
XX transposable element. The transposable elements of the invention are able
XX to introduce in-frame insertions throughout the chromosome of an
XX intracellular bacterial pathogen. This system "tags" the bacterial gene
XX and resulting protein, allowing the identification of proteins
XX secreted across the membranes of the eukaryotic cell infected by the
XX bacterium. In one embodiment, the transposable elements contain an
XX antibiotic resistance cassette, two minimal loxp recombination sites, an
XX MHC class I or class II epitope, and flanking insertion ends. A
XX transposase, such as the Cre recombinase protein, is expressed in trans
XX from a plasmid, or can be included in the transposable element. The Cre
XX recombinase loops out the intervening sequences containing the antibiotic
XX resistance cassette. When the transposable element inserts within a gene,
XX the resolved insertion places the MHC class I or class II epitope in
XX frame with the gene. The transposable elements of the invention are
XX useful for detecting an antigenic epitope of an intracellular bacterial
XX pathogen, such as Salmonella sp., Mycobacterium tuberculosis and Listeria
XX monocytoenes. Certain embodiments of the technology, termed
XX "disseminated insertions of class-I epitopes" (DICE-I; DICE-II for
XX class II epitopes) allow the rapid and accurate identification of
XX proteins involved in bacterial pathogenesis so that such proteins can
XX be used as vaccine and drug targets. Carrier vaccines may be generated
XX by infecting bacteria with a transposable element of the invention
XX which additionally comprises an antigen associated with a disease,
XX preferably cancer or a viral or bacterial disease, operably linked to the
XX MHC epitope DNA of the transposable element. The present sequence
XX represents ovalbumin MHC class I epitope-containing fusion protein
XX encoded by a Tn5-DICE transposable element.
XX
XX Sequence 49 AA:
XX
XX Query Match 100.0%; Score 38; DB 22; Length 49;
XX Best Local Similarity 100.0%; Pred. No. 0.8;
XX Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 1 SIINFEKL 8
XX |||||
XX DB 12 SIINFEKL 19
XX
XX RESULT 110
XX AAB48954
XX ID AAB48954 standard; Protein; 57 AA.
XX

```

```

AC AAB48954;
XX
XX 27-MAR-2001 (first entry)
XX
XX DICE-I ovalbumin MHC class I epitope-containing fusion protein.
XX
XX Transposable element; MHC epitope; major histocompatibility complex;
XX intracellular bacterial pathogen; loxp site; Cre recombinase;
XX insertion end; in-frame fusion; detection; antigen;
XX disseminated insertions of class-I epitopes; DICE-I; transposon Tn5;
XX ovalbumin MHC class I epitope.
XX
XX Synthetic.
XX
XX WO200071158-A1.
XX
XX 30-NOV-2000.
XX
XX 26-MAY-2000; 2000WO-US14687.
XX
XX 26-MAY-1999; 99US-0136210.
XX
XX (UYOR-) UNIV OREGON HEALTH SCI.
XX
XX Heffron FL, Parker DC, Ellefson DD;
XX
XX MPI; 2001-031967/04.
XX
XX N-PSDB; AAC91698.
XX
XX Transposable element for detecting an antigenic epitope of a pathogen,
XX comprising 5' and 3' recombining sites, nucleic acid sequences encoding
XX a selectable marker and major histocompatibility complex (MHC) epitope,
XX and an insertion end -
XX
XX Example 2; Fig 12; 63pp; English.
XX
XX The invention relates to a novel transposable element comprising DNA
XX encoding a selectable marker (e.g., antibiotic resistance) located
XX between a 5' recombining site and a 3' recombining site (e.g., loxp
XX sites); DNA encoding an MHC (major histocompatibility complex) epitope
XX either 5' of the 5' recombining site or 3' of the 3' recombining site;
XX and insertion ends comprising an inverted repeat sequence at the 5' and
XX 3' ends of the transposable element sufficient for integration of the
XX transposable element. The transposable elements of the invention are able
XX to introduce in-frame insertions throughout the chromosome of an
XX intracellular bacterial pathogen. This system "tags" the bacterial gene
XX and resulting protein, allowing the identification of proteins
XX secreted across the membranes of the eukaryotic cell infected by the
XX bacterium. In one embodiment, the transposable elements contain an
XX antibiotic resistance cassette, two minimal loxp recombination sites, an
XX MHC class I or class II epitope, and flanking insertion ends. A
XX transposase, such as the Cre recombinase protein, is expressed in trans
XX from a plasmid, or can be included in the transposable element. The Cre
XX recombinase loops out the intervening sequences containing the antibiotic
XX resistance cassette. When the transposable element inserts within a gene,
XX the resolved insertion places the MHC class I or class II epitope in
XX frame with the gene. The transposable elements of the invention are
XX useful for detecting an antigenic epitope of an intracellular bacterial
XX pathogen, such as Salmonella sp., Mycobacterium tuberculosis and Listeria
XX monocytoenes. Certain embodiments of the technology, termed
XX "disseminated insertions of class-I epitopes" (DICE-I; DICE-II for
XX class II epitopes) allow the rapid and accurate identification of
XX proteins involved in bacterial pathogenesis so that such proteins can
XX be used as vaccine and drug targets. Carrier vaccines may be generated
XX by infecting bacteria with a transposable element of the invention
XX which additionally comprises an antigen associated with a disease,
XX preferably cancer or a viral or bacterial disease, operably linked to the
XX MHC epitope DNA of the transposable element. The present sequence
XX represents an ovalbumin MHC class I epitope-containing fusion protein
XX encoded by the resolved sequence of a DICE-I transposable element.
XX
XX Sequence 57 AA:
XX

```


Query Match 100.0%; Score 38; DB 22; Length 57;
 Best Local Similarity 100.0%; Pred. No. 0.93;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEKL 8
 DB 10 SIINFEKL 17

RESULT 111

AAE13458
 ID AAE13458 standard; Protein; 100 AA.

XX AAE13458;

DT 12-FEB-2002 (first entry)

DE Chicken ovalbumin derived protein domain #1.

XX Antigenic complex; epitope; heat shock protein; HSP; tether; javelin;
 KW major histocompatibility complex; MHC; therapy; immune response;
 XX malignancy; chicken.

OS Gallus gallus.

XX Key Location/Qualifiers

FT Region 1..8 /note= "Javelin sequence"

FT Region 66..73 /note= "MHC class I epitope"

FT Region 74..89 /note= "MHC class II epitope"

PN WO200179259-A1.

PD 25-OCT-2001.

PF 17-APR-2001; 2001WO-US12567.

PR 17-APR-2000; 2000US-197462P.

XX (ROTH/) ROTHMAN J E.

PA (MAYH/) MAYHEW M.

PA (HOEM/) HOE M.

PI Rothman JE, Mayhew M, Hoe M;

XX WPI; 2002-017594/02.

XX A new antigenic complex comprising epitopes non-covalently joined to a

PT heat shock protein by a molecular tether designated a javelin are

PI useful to treat or prevent infectious disease or malignancy -

XX Disclosure; Page 14; 47pp; English.

XX The present invention relates to an antigenic complex, comprising a
 CC number of epitopes non-covalently joined to a heat shock protein (HSP) by
 CC a tethering molecule referred to as javelin which has affinity for the
 CC HSP under physiological conditions, where the epitopes are covalently
 CC joined to the tethering molecule and one epitope is major
 CC histocompatibility complex class I (MHC) and the other MHC class II. The
 CC antigenic complex is used to induce immune responses directed towards the
 CC treatment or prevention of infectious diseases and malignancies. The
 CC present sequence is chicken ovalbumin derived protein domain.

XX Sequence 100 AA;

Query Match 100.0%; Score 38; DB 23; Length 100;

Best Local Similarity 100.0%; Pred. No. 1.7; Indels 0; Gaps 0;
 Matches 8; Conservative 0; Mismatches 0;

QY 1 SIINFEKL 8
 |||||

DB 66 SIINFEKL 73

RESULT 112

AAE13460
 ID AAE13460 standard; Protein; 100 AA.

XX AAE13460;

DT 12-FEB-2002 (first entry)

DE Chicken ovalbumin derived protein domain #3.

XX Antigenic complex; epitope; heat shock protein; HSP; tether; javelin;
 KW major histocompatibility complex; MHC; therapy; immune response;
 XX malignancy; chicken.

OS Gallus gallus.

XX Key Location/Qualifiers

FT Region 58..65 /note= "MHC class I epitope"

FT Region 66..81 /note= "MHC class II epitope"

FT Region 93..100 /note= "Javelin sequence"

PN WO200179259-A1.

PD 25-OCT-2001.

PF 17-APR-2001; 2001WO-US12567.

PR 17-APR-2000; 2000US-197462P.

XX (ROTH/) ROTHMAN J E.

PA (MAYH/) MAYHEW M.

PA (HOEM/) HOE M.

PI Rothman JE, Mayhew M, Hoe M;

XX WPI; 2002-017594/02.

XX A new antigenic complex comprising epitopes non-covalently joined to a

PT heat shock protein by a molecular tether designated a javelin are

PI useful to treat or prevent infectious disease or malignancy -

XX Disclosure; Page 14; 47pp; English.

XX The present invention relates to an antigenic complex, comprising a
 CC number of epitopes non-covalently joined to a heat shock protein (HSP) by
 CC a tethering molecule referred to as javelin which has affinity for the
 CC HSP under physiological conditions, where the epitopes are covalently
 CC joined to the tethering molecule and one epitope is major
 CC histocompatibility complex class I (MHC) and the other MHC class II. The
 CC antigenic complex is used to induce immune responses directed towards the
 CC treatment or prevention of infectious diseases and malignancies. The
 CC present sequence is chicken ovalbumin derived protein domain.

XX Sequence 100 AA;

Query Match 100.0%; Score 38; DB 23; Length 100;

Best Local Similarity 100.0%; Pred. No. 1.7; Indels 0; Gaps 0;
 Matches 8; Conservative 0; Mismatches 0;

QY 1 SIINFEKL 8
 |||||

DB 58 SIINFEKL 65
 |||||
 RESULT 113
 AAE13459
 ID AAE13459 standard; Protein; 103 AA.

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XX AAE13459;
AC
XX 12-FEB-2002 (first entry)
DT
XX Chicken ovalbumin derived protein domain #2.
DE
XX Antigenic complex; epitope; heat shock protein; HSP; tether; javelin;
KM major histocompatibility complex; MHC; therapy; immune response;
KM malignancy; chicken.
XX Gallus gallus.
OS
XX Key Location/Qualifiers
FH Region 1..8
FT Region /note= "Javelin sequence"
FT Region 9..11
FT Region /note= "Linker"
FT Region 69..76
FT Region /note= "MHC class I epitope"
FT Region 77..92
FT Region /note= "MHC class II epitope"
XX WO200179259-A1.
XX 25-OCT-2001.
XX 17-APR-2001; 2001WO-US12567.
XX 17-APR-2000; 2000US-197462P.
XX (ROTH/) ROTHMAN J E.
XX (MAYH/) MAYHEW M.
XX (HOEM/) HOE M.
XX Rothman JE, Mayhew M, Hoe M;
XX WPI; 2002-017594/02.
XX A new antigenic complex comprising epitopes non-covalently joined to a
PT heat shock protein by a molecular tether designated a javelin are
PT useful to treat or prevent infectious disease or malignancy -
XX Disclosure; Page 14; 47pp; English.
XX The present invention relates to an antigenic complex, comprising a
CC number of epitopes non-covalently joined to a heat shock protein (HSP) by
CC a tethering molecule referred to as javelin which has affinity for the
CC HSP under physiological conditions, where the epitopes are covalently
CC joined to the tethering molecule and one epitope is major
CC histocompatibility complex class I (MHC) and the other MHC class II. The
CC antigenic complex is used to induce immune responses directed towards the
CC treatment or prevention of infectious diseases and malignancies. The
CC present sequence is chicken ovalbumin derived protein domain.
XX Sequence 103 AA;
SQ
QY 1 STINFEKL 8
DB 69 STINFEKL 76
Query Match 100.0%; Score 38; DB 23; Length 103;
Best Local Similarity 100.0%; Pred. No. 1.7;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
RESULT 114
AAE13461
ID AAE13461 standard; Protein; 103 AA.
AC AAE13461;
XX 12-FEB-2002 (first entry)
DT

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XX Chicken ovalbumin derived protein domain #4.
DE
XX Antigenic complex; epitope; heat shock protein; HSP; tether; javelin;
KM major histocompatibility complex; MHC; therapy; immune response;
KM malignancy; chicken.
XX Gallus gallus.
OS
XX Key Location/Qualifiers
FH Region 58..65
FT Region /note= "MHC class I epitope"
FT Region 66..81
FT Region /note= "MHC class II epitope"
FT Region 93..95
FT Region /note= "Linker sequence"
FT Region 96..103
FT Region /note= "Javelin sequence"
XX WO200179259-A1.
XX 25-OCT-2001.
XX 17-APR-2001; 2001WO-US12567.
XX 17-APR-2000; 2000US-197462P.
XX (ROTH/) ROTHMAN J E.
XX (MAYH/) MAYHEW M.
XX (HOEM/) HOE M.
XX Rothman JE, Mayhew M, Hoe M;
XX WPI; 2002-017594/02.
XX A new antigenic complex comprising epitopes non-covalently joined to a
PT heat shock protein by a molecular tether designated a javelin are
PT useful to treat or prevent infectious disease or malignancy -
XX Disclosure; Page 14; 47pp; English.
XX The present invention relates to an antigenic complex, comprising a
CC number of epitopes non-covalently joined to a heat shock protein (HSP) by
CC a tethering molecule referred to as javelin which has affinity for the
CC HSP under physiological conditions, where the epitopes are covalently
CC joined to the tethering molecule and one epitope is major
CC histocompatibility complex class I (MHC) and the other MHC class II. The
CC antigenic complex is used to induce immune responses directed towards the
CC treatment or prevention of infectious diseases and malignancies. The
CC present sequence is chicken ovalbumin derived protein domain.
XX Sequence 103 AA;
SQ
QY 1 STINFEKL 8
DB 58 STINFEKL 65
Query Match 100.0%; Score 38; DB 23; Length 103;
Best Local Similarity 100.0%; Pred. No. 1.7;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
RESULT 115
AAR89966
ID AAR89966 standard; Protein; 106 AA.
AC AAR89966;
XX 25-MAR-2003 (updated)
DT 12-SEP-1996 (first entry)
XX Polypeptide sequence.
DE
XX

```

Accession	Key	Location/Qualifiers
XX		
OS	Synthetic.	
KW		Polytope/vaccinia, cytotoxic T lymphocyte; CTL, mouse; epitope; vaccines
KM		major histocompatibility complex; pathogen; HLA diversity; avipox virus;
KM		bacterial vector; rhadovirus vector; ISCOM; influenza nuclear protein;
KM		ovalbumin; cytomegalovirus; adenovirus; sendai virus; P.Berghel; MHC;
KW		circumsporozoite protein; influenza NS1; choriomeningitis virus.
XX		
XX	Key	Location/Qualifiers
FT	Region	4..12
FT		/note= "Cytotoxic T lymphocyte (CTL) epitope #1, isolated from influenza nuclear protein (residues 366-374) "
FT	Region	13..20
FT		/note= "CTL epitope #2, isolated from ovalbumin (residues 257-264) "
FT	Region	21..29
FT		/note= "CTL epitope #3, isolated from influenza nuclear protein (residues 147-155) "
FT	Region	30..37
FT		/note= "CTL epitope #4, isolated from influenza nuclear protein (residues 50-58) "
FT	Region	38..46
FT		/note= "CTL epitope #5, isolated from murine cytomegalovirus pp89 (residues 168-176) "
FT	Region	48..58
FT		/note= "CTL epitope #6, isolated from adenovirus 5 E1A (residues 234-243) "
FT	Region	59..67
FT		/note= "CTL epitope #7, isolated from sendai virus nuclear protein (residues 324-332) "
FT	Region	68..76
FT		/note= "CTL epitope #8, isolated from P.Berghel circumsporozoite protein (residues 249-257) "
FT	Region	77..85
FT		/note= "CTL epitope #9, isolated from influenza NS1 (residues 152-160) "
FT	Misc-difference	78
FT		/note= "encoded by GAC" "
FT	Region	86..94
FT		/note= "CTL epitope #10, isolated from lymphocytic choriomeningitis virus nuclear protein (residues 118-126) "
FT	Region	97..106
FT		/note= "monoclonal antibody epitope" "
FT	Misc-difference	105
FT		/note= "encoded by AGA" "
XX		
PN		W09603144-A1.
XX		
PD		08-FEB-1996.
XX		
PF	27-JUL-1995;	95WO-AU00461.
XX		
PR	08-FEB-1995;	95AU-0001009.
PR	27-JUL-1994;	94AU-0007079.
XX		
PA	(COUN-) COUNCIL QUEENSLAND INST MEDICAL RES.	
PA	(CSIR) COMMONWEALTH SCI & IND RES ORG.	
PA	(UYME) UNIV MELBOURNE.	
PA	(HALL-) HALL INST MEDICAL RES WALTER & ELIZA.	
PA	(BIOT-) BIOTECH AUSTRALIA PTY LTD.	
PA	(CSLC-) CSL LTD.	
XX		
PI	Burrows SR, Coupar BEH, Khanna R, Moss DJ, Sunbrier A;	
PI	Thomson SA;	
XX		
DR	WPI; 1996-116788/12.	
DR	N-PSDB; AAT12413.	
XX		
PT	New poly:epitope cytotoxic T lymphocyte vaccines - comprising a recombinant protein including CTL epitope(s) from pathogens, free of natural flanking sequences	

XX	Claim 5; Fig 5; 46pp; English.
PS	
CC	This sequence represents a polypeptide encoded by a DNA insert of a
CC	recombinant vaccinia virus of the invention. This sequence contains 10
CC	murine cytotoxic T lymphocyte (CTL) epitopes. Each of the epitopes in
CC	this sequence are capable of producing a primary CTL response in mice
CC	with the appropriate major histocompatibility complex (MHC) allele. This
CC	sequence (and the DNA encoding it) can be used in vaccines against
CC	multiple epitopes derived from several different pathogens. The vaccine
CC	could alternatively contain a large number of epitopes from one pathogen
CC	so that MHA diversity of the target population is covered. The vaccines
CC	can be delivered by vaccinia virus, avipox virus, bacterial or
CC	rhabdovirus vectors, or by virus-like particles. The proteins are
CC	preferably administered with ISCOMs when they are delivered directly.
CC	The advantage with these vaccines is that they provide a more diverse
CC	immune response.
CC	(Updated on 25-MAR-2003 to correct PA field.)
SQ	
Sequence	106 AA;
OY	
Query Match	100.0%; Score 38; DB 17; Length 106;
Best Local Similarity:	100.0%; Pred. No. 1.8;
Matches	8; Conservative 0; Mismatches 0; Indels 0; Gaps 0.
Db	13 SIINFEXL 20
OY	1 SIINFEXL 8
ID	AAE13462 standard; Protein; 108 AA.
AC	AAE13462;
DT	12-FEB-2002 (first entry)
DE	Chicken ovalbumin derived protein domain #5.
KW	Antigenic complex; epitope; heat shock protein; HSP; tether; javelin; major histocompatibility complex; MHC; therapy; immune response; malignancy; chicken.
KX	Gallus gallus.
OS	
FH	Key Location/Qualifiers
FT	Region 1..8 /note= "Javelin sequence"
FT	Region 66..73 /note= "MHC class I epitope"
FT	Region 74..89 /note= "MHC class II epitope"
FT	Region 101..108 /note= "Javelin sequence"
PN	WO200179259-A1.
PD	25-OCT-2001.
PF	17-APR-2001; 2001WO-US12567.
PR	17-APR-2000; 2000US-197462P.
PA	(ROTH/) ROTHMAN J E. (MAIH/) MAYHEW M. (HOEM/) HOE M.
PI	Rothman JE, Mayhew M, Hoe M;
DR	WPI; 2002-017594/02.
PT	A new antigenic complex comprising epitopes non-covalently joined to a

PT heat shock protein by a molecular tether designated a javelin are
 PT useful to treat or prevent infectious disease or malignancy -
 XX
 XX
 PS Disclosure; Page 15; 47pp; English.
 CC The present invention relates to an antigenic complex, comprising a
 CC number of epitopes non-covalently joined to a heat shock protein (HSP) by
 CC a tethering molecule referred to as javelin which has affinity for the
 CC HSP under physiological conditions, where the epitopes are covalently
 CC joined to the tethering molecule and one epitope is major
 CC histocompatibility complex class I (MHC) and the other MHC class II. The
 CC antigenic complex is used to induce immune responses directed towards the
 CC treatment or prevention of infectious diseases and malignancies. The
 CC present sequence is chicken ovalbumin derived protein domain.
 CC
 SQ Sequence 108 AA;
 Query Match 100.0%; Score 38; DB 23; Length 108;
 Best Local Similarity 100.0%; Pred. No. 1.8;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEXL 8
 |||||
 DB 66 SIINFEXL 73
 RESULT 117
 AAE13463
 ID AAE13463 standard; Protein; 111 AA.
 AC AAE13463;
 XX
 DT 12-FEB-2002 (first entry)
 DE Chicken ovalbumin derived protein domain #6.
 KW Antigenic complex; epitope; heat shock protein; HSP; tether; javelin;
 KW major histocompatibility complex; MHC; therapy; immune response;
 KW malignancy; chicken.
 XX
 OS Gallus gallus.
 FH Key Location/Qualifiers
 FT Region 1..8
 FT /note= "Javelin sequence"
 FT Region 9..11
 FT /note= "Linker"
 FT Region 69..76
 FT /note= "MHC class I epitope"
 FT Region 77..92
 FT /note= "MHC class II epitope"
 FT Region 104..111
 FT /note= "Javelin sequence"
 PN WO200179259-A1.
 PD 25-OCT-2001.
 PF 17-APR-2001; 2001WO-US12567.
 PR 17-APR-2000; 2000US-197462P.
 XX
 PA (ROTH/) ROTHMAN J E.
 PA (MATH/) MAYHEW M.
 PA (HOEW/) HOE M.
 PI Rothman JE, Mayhew M, Hoe M;
 DR MPI; 2002-017594/02.
 A new antigenic complex comprising epitopes non-covalently joined to a
 PT heat shock protein by a molecular tether designated a javelin are
 PT useful to treat or prevent infectious disease or malignancy -

XX
 PS Disclosure; Page 15; 47pp; English.
 CC The present invention relates to an antigenic complex, comprising a
 CC number of epitopes non-covalently joined to a heat shock protein (HSP) by
 CC a tethering molecule referred to as javelin which has affinity for the
 CC HSP under physiological conditions, where the epitopes are covalently
 CC joined to the tethering molecule and one epitope is major
 CC histocompatibility complex class I (MHC) and the other MHC class II. The
 CC antigenic complex is used to induce immune responses directed towards the
 CC treatment or prevention of infectious diseases and malignancies. The
 CC present sequence is chicken ovalbumin derived protein domain.
 CC
 SQ Sequence 111 AA;
 Query Match 100.0%; Score 38; DB 23; Length 111;
 Best Local Similarity 100.0%; Pred. No. 1.9;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 SIINFEXL 8
 |||||
 DB 69 SIINFEXL 76
 RESULT 118
 AAY52575
 ID AAY52575 standard; Protein; 132 AA.
 AC AAY52575;
 XX
 DT 28-FEB-2000 (first entry)
 DE Amino acid sequence of AOS minigene insert of expression vector pMIN.0.
 KW Chimeric; pan DR epitope; expression vector;
 KW promoter; major histocompatibility complex; MHC; targeting; peptide;
 KW epitope; antigen; presentation; class I; cytosolic pathway;
 KW endoplasmic reticulum; class II; extracellular antigen;
 KW endocytic pathway; helper T lymphocyte; HTL; universal epitope;
 KW cytotoxic T lymphocyte; CTL; immune response; immunogenicity; assay;
 KW vaccine; immunity; infection; pathogen; virus; HIV; HBV; HCV;
 KW hepatitis B; hepatitis C; bacterium; protozoan; tumour cell;
 KW autoimmune disease; activation; antiviral; antimetastatic;
 KW immunoprotective; minigene.
 XX
 OS Synthetic.
 PN WO958658-A2.
 PD 18-NOV-1999.
 PF 13-MAY-1999; 99WO-US10646.
 PR 13-MAY-1998; 98US-0078904.
 PR 15-MAY-1998; 98US-0085751.
 XX
 PA (EPIM-) EPIMUNE INC.
 PI Fikes JD, Hermanson GG, Sette A, Ishioka GY, Livingston B;
 PI Chesnut RW;
 DR MPI; 2000-039103/03.
 DR N-PSDB; AA238634.
 PT Expression vectors encoding major histocompatibility targeting
 PT sequence, used as, e.g. tumor vaccines -
 XX
 PS Example 1; Fig 20; 130pp; English.
 CC This sequence represents the amino acid sequence of the AOS minigene
 CC insert of the expression vector pMIN.0 (AA238634). This insert encodes
 CC several MHC class I epitopes, and also the universal MHC
 CC class II (helper T) epitope, pan DR epitope (PADRE), and was used

in an exemplification of the present invention. The invention relates to a novel expression vector comprising a promoter operably linked to a fusion gene encoding a major histocompatibility complex (MHC) targeting sequence, and two or more heterologous peptide epitopes. The MHC targeting sequence may be a class I targeting sequence, which directs an MHC class I epitope to a cytosolic pathway or to the endoplasmic reticulum, or an MHC class II targeting sequence, which directs extracellular antigens to enter the endocytic pathway to be processed into antigen peptides for presentation on MHC class II molecules. The heterologous epitopes may comprise either helper T lymphocyte (HTL) epitopes, or a cytotoxic T lymphocyte (CTL) epitope and a universal HTL epitope, such as a pan DR epitope (PADRE). The vectors are useful for stimulating an immune response in vivo, as well as for use in assaying the human immunogenicity of a human T cell peptide epitope in vivo in a non-human mammal. They provide a nucleic acid vaccine for enhancing immunity against infectious pathogens, such as viruses (e.g., HIV, hepatitis B (HBV) and hepatitis C (HCV)) bacteria, protozoa (e.g., Plasmodium falciparum, the cause of malaria) and also tumor cells and autoimmune diseases. Universal MHC class II epitopes are advantageously combined with other MHC class I and class II epitopes to increase the number of cells that are activated in response to a given antigen and provide a broader population coverage of MHC-reactive alleles.

SQ Sequence 132 AA;
Query Match 100.0%; Score 38; DB 21; Length 132;
Best Local Similarity 100.0%; Pred. No. 2.2;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEKL 8
Db 75 STINFEKL 82

RESULT 119

AAE13435
ID AAE13435 standard; Protein; 386 AA.

AC AAE13435;

DT 12-FEB-2002 (first entry)

DE Chicken ovalbumin containing plurality of epitopes.

XX Antigenic complex; epitope; heat shock protein; HSP; tether; javelin;
XX major histocompatibility complex; MHC; therapy; immune response;
XX malignancy; chicken.

OS Gallus gallus.

XX Key Location/Qualifiers

FT Domain 200..291

FT Region 258..265

FT Region /note="MHC class I epitope"

FT Region 266..281

FT Region /note="MHC class II epitope"

PN WO200179259-A1.

XX 25-OCT-2001.

PD 17-APR-2001; 2001WO-US12567.

PF 17-APR-2000; 2000US-197462P.

XX (ROTH/) ROTHMAN J E.

PA (MAYH/) MAYHEW M.

PA (HOEM/) HOE M.

XX Rothman JE, Mayhew M, Hoe M;

XX WPI; 2002-017594/02.

DR N-PSDB; AAD22407.

XX A new antigenic complex comprising epitopes non-covalently joined to a
PT heat shock protein by a molecular tether designated a javelin are
PT useful to treat or prevent infectious disease or malignancy -

PS Example; Fig 2; 47pp; English.

XX The present invention relates to an antigenic complex, comprising a
XX number of epitopes non-covalently joined to a heat shock protein (HSP) by
CC a tethering molecule referred to as javelin which has affinity for the
CC HSP under physiological conditions, where the epitopes are covalently
CC joined to the tethering molecule and one epitope is major
CC histocompatibility complex class I (MHC) and the other MHC class II. The
CC antigenic complex is used to induce immune responses directed towards the
CC treatment or prevention of infectious diseases and malignancies. The
CC present sequence is chicken ovalbumin containing plurality of epitopes.

SQ Sequence 386 AA;

QY Query Match 100.0%; Score 38; DB 23; Length 386;
Db Best Local Similarity 100.0%; Pred. No. 6.8;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 STINFEKL 8
Db 258 STINFEKL 265

RESULT 120

AAAB31545
ID AAAB31545 standard; Protein; 409 AA.

AC AAAB31545;

DT 30-APR-2001 (first entry)

DE Amino acid sequence of chicken ovalbumin.

XX Heat shock protein; Hsp; Th1 response; Th1 cell; CD4+ T lymphocyte cell;

XX lymphocyte; Hsp65; Hsp40; Hsp10; Hsp60; Hsp71; microbial pathogen;

XX ovalbumin.

XX Gallus sp.

OS WO200104344-A2.

XX 18-JAN-2001.

PD 10-JUL-2000; 2000WO-US18828.

PF 08-JUL-1999; 99US-0143757.

XX (STRE-) STRESSGEN BIOTECHNOLOGIES CORP.

XX Siegel M, Chu NR, Mizzen LA;

XX WPI; 2001-138361/14.

XX N-PSDB; AAF25127.

PT Screening for compounds that stimulate Th1-like responses in CD4+ T

PT lymphocyte cells -

PS Example 8; Fig 6; 88pp; English.

XX The present sequence represents an ovalbumin protein. Ovalbumin was fused
CC to a heat shock protein (HSP), and used used in the method of the
CC invention. The specification describes a method of determining whether
CC a compound stimulates a Th1-like response. Th1 cells are a subset of
CC CD4+ T lymphocyte cells. The method comprises contacting naive
CC lymphocytes in vitro with a fusion protein comprising at least a
CC fragment of Hsp, and then detecting the Th1-like response exhibited by
CC the cell sample. The proteins which may be used in the method of the
CC invention are Hsp65, Hsp40, Hsp10, Hsp60, and Hsp71. The method may be

CC used to identify compounds that stimulate Th1-like responses in response
CC to microbial pathogens.

XX Sequence 409 AA;

Query Match 100.0%; Score 38; DB 22; Length 409;

Best Local Similarity 100.0%; Pred. No. 7.2;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINEFKL 8
|||||

DB 281 SIINEFKL 288

RESULT 121

ID AAE13112 standard; Protein; 479 AA.

AC AAE13112;

DT 28-JAN-2002 (first entry)

DE Human HER300-IGM-CSF fusion construct comprising OVA-derived peptide.

XX Immunostimulatory fusion protein; IFP; antigen component; therapy;
KW immunostimulatory component; T-cell mediated immune response; DC;
KW dendritic cell; colon cancer; breast carcinoma; ovarian cancer;
KW PAP protein; Ala Arg linker; membrane distal extracellular domain;
KW membrane distal intracellular domain; C-terminal tag; human; GM-CSF;
KW HER-2 protein; granulocyte-macrophage colony stimulating factor;
KW ovalbumin-derived octapeptide; OVA; rat; HER300-IGM-CSF fusion protein.

OS Chimeric - Homo sapiens.

OS Chimeric - Rattus norvegicus.

OS Chimeric - Unidentified.

PN WO200174855-A2.

PD 11-OCT-2001.

PR 30-MAR-2001; 2001WO-US10515.

PA 30-MAR-2000; 2000US-193504P.

PI (DEND-) DENDREON CORP.

PI Laus R, Vidovic D, Graddis T;

DR WPI; 2001-662365/76.

DR N-PSDB; AAD21568.

PT An immunostimulatory fusion protein comprising the intracellular domain
PT of HER-2 and an antigen elicits an immune response to the antigen and
PT is useful for the treatment of associated cancer associated -

PS Example 1; Page 27; 59pp; English.

XX The invention relates to immunostimulatory fusion proteins (IFP) and
CC nucleic acid molecules encoding such proteins. The IFPs comprise a
CC polypeptide antigen component and an immunostimulatory component derived
CC from the intracellular domain of HER-2 protein which is effective to
CC elicit a protective dendritic cell (DC)-induced T-cell mediated cellular
CC immune response to the antigen. IFP or superactivated dendritic cells
CC are used to treat cancer e.g. breast carcinoma, ovarian and colon cancer
CC associated with a particularly antigen. The present sequence is HER300
CC IGM-CSF fusion protein construct which comprises human PAP
CC signal sequence, mature PAP protein, an Ala Arg linker, human HER-2
CC signal sequence, mature HER-2 membrane distal extracellular domain,
CC an Ala linker, an ovalbumin (OVA)-derived immunodominant octapeptide,
CC an Ala linker, a HER-2 membrane distal intracellular domain, a mature
CC rat granulocyte-macrophage colony stimulating factor (GM-CSF) sequence
CC and a C-terminal tag.

SQL Sequence 479 AA;

Query Match 100.0%; Score 38; DB 22; Length 479;

Best Local Similarity 100.0%; Pred. No. 8.5;
Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINEFKL 8
|||||

DB 331 SIINEFKL 338

RESULT 122

ID AAU99725 standard; Protein; 541 AA.

AC AAU99725;

DT 07-OCT-2002 (first entry)

DE Yeast/mouse SS-OVA-Kb/beta2m-c-myc-AGA2 fusion protein.

XX Mutant major histocompatibility complex class I chimeric protein; MHC;
KW lymphocyte; T-cell receptor; tissue sample; biopsy material; pathogen;
KW bodily fluid; T lymphocyte; neoplastic cell; tumour cell; MHC antigen;
KW virus; protozoan; bacteria; fungi; nematode; immune response; activator;
KW enhancer; T cell activator; mouse; recombinant yeast cell; Kb; OVA;
KW beta2m; dev8; AGA2; SIYK; fusion protein.

OS Chimeric - Mus sp.

OS Chimeric - Saccharomyces cerevisiae.

OS Synthetic.

PN WO200246399-A2.

PD 13-JUN-2002.

PR 10-DEC-2001; 2001WO-US47817.

PA 08-DEC-2000; 2000US-254495P.

PI (UNIT) UNIV ILLINOIS FOUNO.

PI Kranz DM, Brophy S;

DR WPI; 2002-527916/56.

DR N-PSDB; ABE87870.

PT New isolated mutant major histocompatibility complex class I chimeric
PT protein displayed on surfaces of recombinant yeast cells, has improved
PT stability, and is useful for activating immune response -

PS Example 7; Page 38-39; 96pp; English.

XX The present invention relates to a new mutant major histocompatibility
CC complex (MHC) class I chimeric protein. The protein of the invention
CC comprises a portion mediating binding to surfaces of recombinant yeast
CC cells and a portion comprising peptide binding region of MHC class I
CC protein, where the invention is improved in stability as compared with
CC MHC class I chimeric protein which is not a mutant chimeric protein.
CC The protein, further comprising a detectable label, is useful for
CC detecting a lymphocyte having a T-cell receptor protein in a biological
CC sample such as cells, tissue sample, biopsy material or bodily fluids.
CC The method is useful for detecting a T lymphocyte that is specific for
CC a neoplastic cell, a tumour cell, a virus-infected cell, a protozoan-
CC infected cell, a bacterium-infected cell or a fungus-infected cell. The
CC protein of the invention can be used to directly activate T cells, in
CC order to identify/screen for peptide-MHC antigens. The protein is also
CC useful in activating T cells that participate in the removal of target
CC cells including neoplastic cells and cells infected with pathogenic
CC agents including viruses, protozoans, bacteria, fungi or nematodes.
CC The invention is improved in stability as compared with MHC class I
CC protein which is not a mutant chimeric protein. The present amino acid
CC sequence represents a chimeric MHC protein that is encoded by a

CC yeast/mouse fusion gene, as described above.
 XX Sequence 541 AA;

Query Match 100.0%; Score 38; DB 23; Length 541;
 Best Local Similarity 100.0%; Pred. No. 9.7;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEKL 8
 |||||
 DB 34 SIINFEKL 41

RESULT 123
 AAE13110
 ID AAE13110 standard; Protein; 564 AA.

XX AAE13110;
 XX
 DT 28-JAN-2002 (first entry)

DE Human HER500 fusion protein construct comprising OVA-derived octapeptide.

XX Immunostimulatory fusion protein; IFP; antigen component; therapy;
 KW immunostimulatory component; T-cell mediated immune response; DC;
 KW dendritic cell; colon cancer; breast carcinoma; ovarian cancer;
 KW PAP protein; Ala Arg linker; membrane distal extracellular domain;
 KW membrane distal intracellular domain; C-terminal tag; human; OVA;
 KW HER-2 protein; ovalbumin-derived octapeptide; HER500 fusion protein.

XX OS Chimeric - Homo sapiens.
 OS Chimeric - unidentified.

PN WO200174855-A2.

XX 11-OCT-2001.

XX 30-MAR-2001; 2001WO-US10515.

XX 30-MAR-2000; 2000US-193504P.

XX (DEND-) DENDREON CORP.

PI Laus R, Vidovic D, Graddis T;

XX WPI; 2001-662965/76.

DR N-PSDB; AAD21566.

PT An immunostimulatory fusion protein comprising the intracellular domain
 of HER-2 and an antigen elicits an immune response to the antigen and
 is useful for the treatment of associated cancer associated -
 XX
 PS Claim 7; Page 26; 59pp; English.

CC The invention relates to immunostimulatory fusion proteins (IFP) and
 CC nucleic acid molecules encoding such proteins. The IFPs comprise a
 CC polypeptide antigen component and an immunostimulatory component derived
 CC from the intracellular domain of HER-2 protein which is effective to
 CC elicit a protective dendritic cell (DC)-induced T-cell mediated cellular
 CC immune response to the antigen. IFP or superactivated dendritic cells
 CC are used to treat cancer e.g. breast carcinoma, ovarian and colon cancer
 CC associated with a particularly antigen. The present sequence is HER500
 CC fusion protein construct which comprises human PAP signal
 CC sequence, mature PAP protein, an Ala Arg linker, human HER-2 signal
 CC sequence, mature HER-2 membrane distal extracellular domain, an
 CC Ala linker, an ovalbumin (OVA)-derived immunodominant octapeptide,
 CC HER-2 membrane distal intracellular domain and a C-terminal tag.

XX Sequence 564 AA;

Query Match 100.0%; Score 38; DB 22; Length 564;
 Best Local Similarity 100.0%; Pred. No. 10;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 SIINFEKL 8
 |||||
 DB 331 SIINFEKL 338

RESULT 124
 AAE13111
 ID AAE13111 standard; Protein; 697 AA.

XX AAE13111;
 XX
 DT 28-JAN-2002 (first entry)

DE Human HER500-rGM-CSF fusion construct comprising OVA-derived peptide.

XX Immunostimulatory fusion protein; IFP; antigen component; therapy;
 KW immunostimulatory component; T-cell mediated immune response; DC;
 KW dendritic cell; colon cancer; breast carcinoma; ovarian cancer;
 KW PAP protein; Ala Arg linker; membrane distal extracellular domain;
 KW membrane distal intracellular domain; C-terminal tag; human; GM-CSF;
 KW HER-2 protein; granulocyte-macrophage colony stimulating factor;
 KW ovalbumin-derived octapeptide; OVA; rat; HER500-rGM-CSF fusion protein.

XX OS Chimeric - Homo sapiens.
 OS Chimeric - Rattus norvegicus.
 OS Chimeric - unidentified.

PN WO200174855-A2.

XX 11-OCT-2001.

XX 30-MAR-2001; 2001WO-US10515.

XX 30-MAR-2000; 2000US-193504P.

XX (DEND-) DENDREON CORP.

PI Laus R, Vidovic D, Graddis T;

XX WPI; 2001-662965/76.

DR N-PSDB; AAD21567.

PT An immunostimulatory fusion protein comprising the intracellular domain
 of HER-2 and an antigen elicits an immune response to the antigen and
 is useful for the treatment of associated cancer associated -
 XX
 PS Claim 7; Page 27; 59pp; English.

CC The invention relates to immunostimulatory fusion proteins (IFP) and
 CC nucleic acid molecules encoding such proteins. The IFPs comprise a
 CC polypeptide antigen component and an immunostimulatory component derived
 CC from the intracellular domain of HER-2 protein which is effective to
 CC elicit a protective dendritic cell (DC)-induced T-cell mediated cellular
 CC immune response to the antigen. IFP or superactivated dendritic cells
 CC are used to treat cancer e.g. breast carcinoma, ovarian and colon cancer
 CC associated with a particularly antigen. The present sequence is HER500
 CC rGM-CSF fusion protein construct which comprises human PAP
 CC signal sequence, mature PAP protein, an Ala Arg linker, human HER-2
 CC signal sequence, mature HER-2 membrane distal extracellular domain,
 CC an Ala linker, an ovalbumin (OVA)-derived immunodominant octapeptide,
 CC HER-2 membrane distal intracellular domain, an Ala Ala linker, a mature
 CC rat granulocyte-macrophage colony stimulating factor (GM-CSF) sequence
 CC and a C-terminal tag.

XX Sequence 697 AA;

Query Match 100.0%; Score 38; DB 22; Length 697;
 Best Local Similarity 100.0%; Pred. No. 13;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 1 SIINFEKL 8
 |||||

Db 331 SIINFEXL 338

RESULT 125

AA031611 AAB31611 standard; Protein; 948 AA.

AC AAB31611;

DT 30-APR-2001 (first entry)

DE Amino acid sequence of Hsp65-ovalbumin fusion protein.

KW Heat shock protein; Hsp; Th1 response; Th1 cell; CD4+ T lymphocyte cell;

KW lymphocyte; Hsp65; Hsp40; Hsp10; Hsp60; Hsp71; microbial pathogen;

OS Synthetic.

OS Mycobacterium bovis.

PN WO200104344-A2.

PD 18-JAN-2001.

PF 10-JUL-2000; 2000WO-US18828.

PR 08-JUL-1999; 99US-0143757.

PA (STRE-) STRESSGEN BIOTECHNOLOGIES CORP.

PI Siegel M, Chu NR, Mizzzen LA;

DR WPI: 2001-138361/14.

DR N-PSDB; AAF25014.

PT Screening for compounds that stimulate Th1-like responses in CD4+ T

PS Example 8; Fig 7A-B; 88pp; English.

CC The present sequence represents a fusion protein comprising a

CC Mycobacterium bovis heat shock protein (Hsp) 65 fused at its C terminal

CC to an ovalbumin protein. The fusion protein is used in the method of the

CC invention. The specification describes a method of determining whether a

CC compound stimulates a Th1-like response. Th1 cells are a subset of CD4+

CC T lymphocyte cells. The method comprises contacting naive lymphocytes

CC in vitro with a fusion protein comprising at least a fragment of Hsp,

CC and then detecting the Th1-like response exhibited by the cell sample.

CC The proteins which may be used in the method of the invention are Hsp65,

CC Hsp40, Hsp10, Hsp60, and Hsp71. The method may be used to identify

CC compounds that stimulate Th1-like responses in response to microbial

CC pathogens.

SQ Sequence 948 AA;

Query Match 100.0%; Score 38; DB 22; Length 948;

Best Local Similarity 100.0%; Pred. No. 17;

Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 SIINFEXL 8

Db 820 SIINFEXL 827

Search completed: January 30, 2004, 07:17:23
Job time : 74 secs

! FINDPATTERNS on geneseqp: * allowing 0 mismatches
! 1 C(R,K){20,20} January 30, 2004 07:26 ..

1 AAW45801 ck: 9500 len: 39 ! Aaw45801 One chain of a bombesin dimer. 6/1

1: C(R,K){20,20}
C(K){20}
CKKKKKKKKKKKKKKKKKKK XGGGQ

1 AAB13780 ck: 7317 len: 21 ! Aab13780 Soluble peptide antigen PK: 11/200

1: C(R,K){20,20}
C(K){20}
CKKKKKKKKKKKKKKKKKKK

1 ABG92659 ck: 5509 len: 58 ! Abg92659 Human DNA-binding protein #85. 11/

1: C(R,K){20,20}
C(K){20}
CKKKKKKKKKKKKKKKKKKK KKK

1 AAU18238 ck: 5509 len: 58 ! Aau18238 Novel human DNA-binding protein #8

1: C(R,K){20,20}
C(K){20}
CKKKKKKKKKKKKKKKKKKK KKK

1 AA003766 ck: 8808 len: 81 ! Aa003766 Human polypeptide SEQ ID NO 17658.

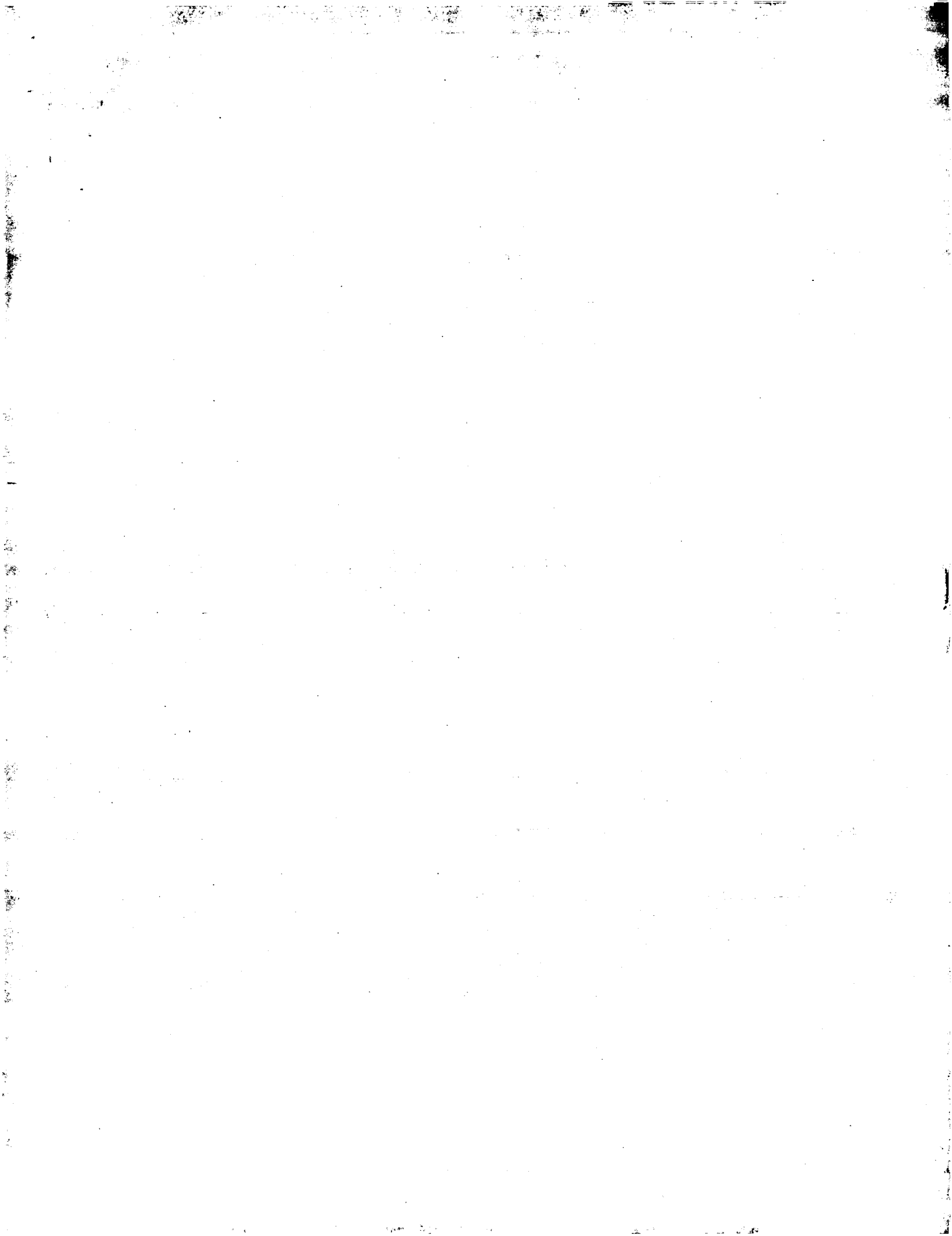
1: C(R,K){20,20}
C(K){20}
LTTTA CKKKKKKKKKKKKKKKKKKK KKKKK

1 AA011210 ck: 863 len: 70 ! Aa011210 Human polypeptide SEQ ID NO 25102.

1: C(R,K){20,20}
C(K){20}
30: IDLCL CKKKKKKKKKKKKKKKKKKK KKKKK

Databases searched:
Geneseq-AA, Release 13.0, Released on 19Jun2003, Formatted on 15Jul2003

Total finds: 6
Total length: 158,726,570
Total sequences: 1,107,863
CPU time: 04:40.14



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!!SEQUENCE LIST 1.0
! FINDPATTERNS on geneseqp:* allowing 0 mismatches
!      1 C(R,K){20,20}      January 30, 2004 07:38 ..
```

```
GENESEQP1990S:AAW45801 ck: 9500 len: 39 finds: 1 | Aaw45801 One chain of a bomb
GENESEQP2000S:AAB13780 ck: 7317 len: 21 finds: 1 | Aab13780 Soluble peptide and
GENESEQP2002S:ABG92659 ck: 5509 len: 58 finds: 1 | Abg92659 Human DNA-binding P
GENESEQP2001S:AAU18238 ck: 5509 len: 58 finds: 1 | Aau18238 Novel human DNA-bir
GENESEQP2001S:AAO03766 ck: 8808 len: 81 finds: 1 | Aao03766 Human polypeptide S
GENESEQP2001S:AAO11210 ck: 863 len: 70 finds: 1 | Aao11210 Human polypeptide S
```

\\End of list

Databases searched:

Geneseq-AA, Release 13.0, Released on 19Jun2003, Formatted on 15Jul2003

```
Total finds: 6
Total length: 158,726,570
Total sequences: 1,107,863
CPU time: 07:02.46
```



```

!!AA_SEQUENCE 1.0
ID_AAW45801 standard; peptide: 39 AA.
XX
AC AAW45801;
XX
DT 25-JUN-1998 (first entry)
XX
DE One chain of a bombesin dimer.
XX
KW Alpha-melanocyte stimulating hormone; alpha-MSH; receptor agonist;
KM alpha-MSH-ANP; bombesin; dimer; bivalent agonist; disulphide bond;
XX G-protein coupled receptor.
OS Synthetic.
XX
FH Key Location/Qualifiers
FT Cross-links 1 /note= "This residue is disulphide bonded to the
FT corresponding Cys residue of an identical chain
FT to form a dimer"
FT Modified-site 28 /note= "Epsilon-aminohexanoic acid"
PN MO9803632-AI.
PD 29-JAN-1998.
PE 23-JUL-1997; 97WO-US12911.
PR 24-JUL-1996; 96US-0686934.
PA (UYYA ) UNIT YALE.
PI Carltchers MD, Lerner MR;
XX WPI; 1998-120757/11.
DR
XX Bivalent agonist of G-protein coupled receptors containing two
PT ligand domains - bonded to molecular backbone, for treatment of
PT hypotension, promotion of skin tanning etc.; also for delivering
PT drugs and gene therapy vectors to selected cells
XX
PS Claim 41; Page 48; 71pp; English.
XX
CC This sequence represents one of two identical chains disulphide bonded
CC to form a bombesin dimer. The invention relates to bivalent agonists,
CC with affinity for at least 1 G-protein coupled receptor (GPCR). The
CC bivalent agonists comprise: (a) two ligand domains (LD), individually
CC agonists or antagonists for GPCR, spaced 40-250 Angstrom apart, and (b)
CC a molecular backbone (MB) covalently bound to LD. The bivalent agonists
CC are useful in human or veterinary medicine as carriers for drugs or gene
CC therapy vectors, allowing these to be endocytosed by GPCR-expressing
CC cells. They can also be used e.g. to treat hypertension (angiotensin-
CC based LD); to increase levels of lutealising hormone (LH), using LD
CC derived from LH-releasing hormone, or to promote skin tanning (LD based
CC on alpha-melanocyte-stimulating hormone, MSH). The bivalent agonists
CC are administered orally, by injection or topically. Typical doses for
CC skin tanning are 1-4000 (especially 30-100) mu mole/kg systemically or
CC the bivalent agonists are administered topically in a composition
CC containing 0.001-10 (especially 1) mM. Where both LD are agonists, the
CC bivalent agonist has a synergistically higher activity than two
CC individual agonist ligands, and where at least 1 is an antagonist the
CC effect is stimulatory. The bivalent agonists are active at lower
CC concentrations than known agonists so should avoid toxicity problems.
S0 Sequence 39 AA;
AAW45801 Length: 39 January 30, 2004 07:48 Type: P Check: 9500 ..
1 CKKKKKKKK KKKKKKKKK KXGGGQGRLG NQNAVGHLM

```

XX	AAB13780;
AC	
XX	10-NOV-2000 (first entry)
DT	
XX	
DE	Soluble peptide antigen pK.
XX	
KM	pK peptide; cytostatic; vaccine; cytotoxic T cell; CTL; immunotherapy;
KW	major histocompatibility complex class I; MHC class I; antigen; tumour;
KM	prostate; breast; multiple myeloma.
OS	
XX	Unidentified.
FN	
XX	WO200035949-A1.
PD	
XX	22-JUN-2000.
PF	
XX	14-DEC-1999; 99WO-US29724.
PR	
XX	14-DEC-1998; 98US-0112324.
PA	(DEND-) DENDREON CORP.
PI	
XX	Laus R, Hakim I, Vidovic D;
DR	WPI; 2000-442365/38.
XX	
PT	Antigens modified by the covalent addition of a peptide that
PT	facilitates entry into antigen presenting cells, useful for producing
PT	compositions for immunizing against tumors and pathogens -
PS	
XX	Claim 2; Page 26; 34pp; English.
CC	The present invention relates to compositions of modified soluble protein
CC	antigens capable of eliciting an enhanced in vivo cytotoxic T cell (CTL)
CC	response i.e. a major histocompatibility complex (MHC) class I molecule
CC	response. The protein antigen is modified by the covalent addition of a
CC	peptide sequence which facilitate entry of the antigen into antigen
CC	presenting cells (APCs). The present sequence is one such peptide
CC	sequence which can be used to modify the soluble antigens. The present
CC	sequence is peptide pK. The modified antigen composition may be used for
CC	immunising against, or creating a tumour e.g. prostate and breast
CC	carcinoma or multiple myeloma, or pathogen in mammals.
SQ	
Sequence	21 AA;
AAB13780	Length: 21 January 30, 2004 07:48 Type: P Check: 7317 ..
1	CKKKKKKKKK KKKKKKKKK K
IIAA_SEQUENCE 1.0	
ID	_ABG92659 standard; Protein; 58 AA.
XX	
XC	ABG92659;
XX	
DT	18-NOV-2002 (first entry)
XX	
DE	Human DNA-binding protein #85.
XX	
KM	Human; DNA-binding protein; B cell immunodeficiency; autoimmune disorder;
KM	severe combined immunodeficiency; rheumatoid arthritis; Crohn's disease;
KM	diabetes mellitus; allergy; asthma; inflammatory condition; thrombosis;
KM	graft-versus-host disease; blood-related disorder; atherosclerosis;
KM	hyperproliferative disease; cancer; renal disorder; arrhythmia;
KM	acute glomerulonephritis; cardiovascular disorder; respiratory disorder;
KM	Gout pasture s syndrome; neurological disorder; Alzheimer's disease;
KM	Parkinson's disease; endocrine disorder; Addison's disease;
KM	reproductive system disorder; endometriosis; infectious disease;
KM	viral infection; bacterial infection; fungal infection; vaccine;
KM	gastrointestinal disorder; multiple sclerosis; gene therapy.
XX	
SS	Homo sapiens.
XX	

```

PN US2002102638-A1.
XX
XX 01-AUG-2002.
XX
PF 17-JAN-2001; 2001US-0764846.
XX
PR 31-JAN-2000; 2000US-179065P.
PR 04-FEB-2000; 2000US-180628P.
PR 28-JUN-2000; 2000US-214886P.
PR 07-JUL-2000; 2000US-216647P.
PR 07-JUL-2000; 2000US-216880P.
PR 11-JUL-2000; 2000US-217487P.
PR 11-JUL-2000; 2000US-217496P.
PR 14-JUL-2000; 2000US-218290P.
PR 26-JUL-2000; 2000US-220963P.
PR 26-JUL-2000; 2000US-220964P.
PR 14-AUG-2000; 2000US-224548P.
PR 14-AUG-2000; 2000US-224519P.
PR 14-AUG-2000; 2000US-225267P.
PR 14-AUG-2000; 2000US-225268P.
PR 14-AUG-2000; 2000US-225270P.
PR 14-AUG-2000; 2000US-225447P.
PR 14-AUG-2000; 2000US-225757P.
PR 14-AUG-2000; 2000US-225758P.
PR 22-AUG-2000; 2000US-226688P.
PR 30-AUG-2000; 2000US-228924P.
PR 01-SEP-2000; 2000US-229287P.
PR 01-SEP-2000; 2000US-229343P.
PR 01-SEP-2000; 2000US-229344P.
PR 01-SEP-2000; 2000US-229345P.
PR 05-SEP-2000; 2000US-229509P.
PR 05-SEP-2000; 2000US-229513P.
PR 08-SEP-2000; 2000US-231413P.
PR 21-SEP-2000; 2000US-234223P.
PR 21-SEP-2000; 2000US-234274P.
PR 25-SEP-2000; 2000US-234997P.
PR 27-SEP-2000; 2000US-235834P.
PR 29-SEP-2000; 2000US-236337P.
PR 29-SEP-2000; 2000US-236367P.
PR 29-SEP-2000; 2000US-236368P.
PR 29-SEP-2000; 2000US-236369P.
PR 29-SEP-2000; 2000US-236370P.
PR 02-OCT-2000; 2000US-236802P.
PR 02-OCT-2000; 2000US-237037P.
PR 02-OCT-2000; 2000US-237038P.
PR 02-OCT-2000; 2000US-237039P.
PR 02-OCT-2000; 2000US-237040P.
PR 13-OCT-2000; 2000US-239935P.
PR 20-OCT-2000; 2000US-240960P.
PR 20-OCT-2000; 2000US-241785P.
PR 20-OCT-2000; 2000US-241809P.
PR 01-NOV-2000; 2000US-244617P.
PR 17-NOV-2000; 2000US-249299P.
PR 08-DEC-2000; 2000US-251856P.
PR 08-DEC-2000; 2000US-251868P.
PR 08-DEC-2000; 2000US-251869P.
XX
XX (ROSE/) ROSEN C A.
XX (RUBE/) RUBEN S M.
XX (BARA/) BARASH S C.
XX
XX Rosen CA, Ruben SM, Barash SC,
XX
XX WPI; 2002-690611/74.
XX N-PSDB; ABS68254.
XX
XX Novel DNA-binding protein useful for diagnosis, prognosis, prevention
XX and treatment of immune, hyperproliferative, respiratory,
XX cardiovascular, reproductive, endocrine, gastrointestinal and
XX neurological disorders
XX
XX Claim 11; SEQ ID No 223; 225bp; English.
XX

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CC The present invention relates to a new DNA-binding protein. The invention
CC is useful in treating, preventing, diagnosing and/or prognosing
CC immunodeficiencies (e.g. B cell immunodeficiencies, severe combined
CC immunodeficiencies), autoimmune disorders (rheumatoid arthritis, multiple
CC sclerosis, diabetes mellitus), allergic reactions and conditions (e.g.
CC asthma), inflammatory conditions, graft-versus-host disease, blood-
CC related disorders (thrombosis, atherosclerosis), hyperproliferative
CC disorders (e.g. cancer), renal disorders (e.g. acute glomerulonephritis),
CC cardiovascular disorders (e.g. arrhythmia), respiratory disorders
CC (COPD/pasture's syndrome), neurological disorders (e.g. Alzheimer's
CC disease, Parkinson's disease), endocrine disorders (e.g. Addison's
CC disease), reproductive system disorders (e.g. endometriosis),
CC infectious diseases (e.g. viral, bacterial or fungal infections) and
CC gastrointestinal disorders (e.g. Crohn's disease). The invention is also
CC useful to stimulate neuronal growth and treat, prevent, and/or diagnose
CC neuronal damage which occurs in certain neuronal disorders or neuro-
CC degenerative conditions. The present amino acid sequence represents a
CC human DNA-binding protein of the invention.
CC Note: The sequence data for this patent did not form part of the
CC printed specification, but was obtained in electronic format directly
CC from USPTO at
CC http://seqdata.uspto.gov/sequence.
XX
XX SQ Sequence 58 AA;
XX
XX ABG92659 Length: 58 January 30, 2004 07:48 Type: P Check: 5509
XX
XX 1 TYLECEHNSL VNSKCLTVVL SRCISVCINK FFFVCKKKKK KKKKKKKKKK
XX
XX 51 KKKKKKKK
XX
XX !:AA SEQUENCE 1.0
XX ID _AAU18238 standard; Protein: 58 AA.
XX
XX AAU18238;
XX
XX 21-NOV-2001 (first entry)
XX
XX DE Novel human DNA-binding protein #85.
XX
XX Human, DNA-binding protein; histone; chromo domain protein;
XX chromatin organisation modifier; Y-box binding protein;
XX DNA organisation; gene transcription; malignant disease;
XX autoimmune disorder; rheumatic disease; genetic abnormality;
XX infectious disease; neurological disorder; gene therapy;
XX immunomodulatory; anti-HIV; anti rheumatic; anti microbial;
XX cytostatic.
XX
XX OS Homo sapiens.
XX
XX WO200155162-A1.
XX
XX 02-AUG-2001.
XX
XX PF 17-JAN-2001; 2001WO-US01305.
XX
XX
XX 31-JAN-2000; 2000US-0179065.
XX 04-FEB-2000; 2000US-0180628.
XX 24-FEB-2000; 2000US-0184664.
XX 02-MAR-2000; 2000US-0186350.
XX 16-MAR-2000; 2000US-0189874.
XX 17-MAR-2000; 2000US-0190076.
XX 18-APR-2000; 2000US-0198123.
XX 19-MAY-2000; 2000US-0205515.
XX 07-JUN-2000; 2000US-0209467.
XX 28-JUN-2000; 2000US-0214886.
XX 30-JUN-2000; 2000US-0215135.
XX 07-JUL-2000; 2000US-0216647.
XX 07-JUL-2000; 2000US-0216880.
XX 11-JUL-2000; 2000US-0217487.
XX 11-JUL-2000; 2000US-0217496.
XX 14-JUL-2000; 2000US-0218290.
XX 26-JUL-2000; 2000US-0220963.
XX

```


CC disease). The polynucleotide sequences of the invention may also be
 CC used in gene therapy. AAU18154-AAU18281 represent novel DNA-binding
 CC proteins.
 CC Note: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 58 AA;

AAU18238 Length: 58 January 30, 2004 07:48 Type: P Check: 5509 ..

1 TYLCHEHNSL VNSKCLTVVL SRCISVCLNK FYFVCKKKKK KKKKKKKKK

51 KKKKKKKK

11AA_SEQUENCE 1.0

ID_AA03766 standard; Protein; 81 AA.

XX AA03766;

XX 06-NOV-2001 (first entry)

XX Human polypeptide SEQ ID NO 17658.

XX Human; cytokine; cell proliferation; cell differentiation; gene therapy;
 KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
 KW tissue growth factor; immunomodulatory; cancer; leukaemia;
 KW nervous system disorders; arthritis; inflammation.

XX Homo sapiens.

XX WO200164835-A2.

XX 07-SEP-2001.

XX 26-FEB-2001; 2001WO-US04927.

XX 28-FEB-2000; 2000US-0515126.

XX 18-MAY-2000; 2000US-0577409.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Drmanac RT;

XX WPI: 2001-514838/56.

XX N-PSDB; AA183637.

XX Isolated nucleic acids and polypeptides, useful for preventing

XX diagnosing and treating e.g. leukaemia, inflammation and immune

XX disorders -

XX Claim 20; SEQ ID NO 17658; 1399pp + Sequence Listing; English.

XX The invention relates to human polynucleotides (AA179941-AA193841) and
 CC the encoded proteins (AA000010-AA013910) that exhibit activity relating to
 CC cytokine, cell proliferation or cell differentiation or which may induce
 CC production of other cytokines in other cell populations. The
 CC polynucleotides and polypeptides are useful in gene therapy, vaccines or
 CC peptide therapy. The polypeptides have various cytokine-like activities,
 CC e.g. stem cell growth factor activity, haematopoiesis regulating
 CC activity, tissue growth factor activity, immunomodulatory activity and
 CC activin/inhibin activity and may be useful in the diagnosis and/or
 CC treatment of cancer, leukaemia, nervous system disorders, arthritis and
 CC inflammation.
 CC Note: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 81 AA;

AA03766 Length: 81 January 30, 2004 07:48 Type: P Check: 8808 ..

1 GUNOTQLRKI LAVSSITHIG XIAVLPYNP NITILNTIY IILTTRACK

51 KKKKKKKKK KKKKKKKKK KKKKKGGG A

11AA_SEQUENCE 1.0

ID_AA011210 standard; Protein; 70 AA.

XX AA011210;

XX 06-NOV-2001 (first entry)

XX Human polypeptide SEQ ID NO 25102.

XX Human; cytokine; cell proliferation; cell differentiation; gene therapy;
 KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;
 KW tissue growth factor; immunomodulatory; cancer; leukaemia;
 KW nervous system disorders; arthritis; inflammation.

XX Homo sapiens.

XX WO200164835-A2.

XX 07-SEP-2001.

XX 26-FEB-2001; 2001WO-US04927.

XX 28-FEB-2000; 2000US-0515126.

XX 18-MAY-2000; 2000US-0577409.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Drmanac RT;

XX WPI: 2001-514838/56.

XX N-PSDB; AA191141.

XX Isolated nucleic acids and polypeptides, useful for preventing
 XX diagnosing and treating e.g. leukaemia, inflammation and immune
 XX disorders -

XX Claim 20; SEQ ID NO 25102; 1399pp + Sequence Listing; English.

XX The invention relates to human polynucleotides (AA179941-AA193841) and
 CC the encoded proteins (AA000010-AA013910) that exhibit activity relating to
 CC cytokine, cell proliferation or cell differentiation or which may induce
 CC production of other cytokines in other cell populations. The
 CC polynucleotides and polypeptides are useful in gene therapy, vaccines or
 CC peptide therapy. The polypeptides have various cytokine-like activities,
 CC e.g. stem cell growth factor activity, haematopoiesis regulating
 CC activity, tissue growth factor activity, immunomodulatory activity and
 CC activin/inhibin activity and may be useful in the diagnosis and/or
 CC treatment of cancer, leukaemia, nervous system disorders, arthritis and
 CC inflammation.
 CC Note: The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 70 AA;

AA011210 Length: 70 January 30, 2004 07:48 Type: P Check: 863 ..

1 YYIHRTVEM CMNXGLKDNV DKXTIDLCLC KKKKKKKKK KKKKKKKKK

51 KKKKKKKKK KKKKPGGGA

! FINDPATTERNS on pir:* allowing 0 mismatches
! 1 C(R,K) {20,20}

January 30, 2004 06:58 ..

Databases searched:

NBRF, Release 76.1, Released on 12May2003, Formatted on 10Jun2003

Total finds: 0
Total length: 96,168,682
Total sequences: 283,308
CPU time: 01:56.42

! FINDPATTERNS on swp:* allowing 0 mismatches
! 1 C(R,K) {20,20} January 30, 2004 06:58 ..

Databases searched:
SWISS-PROT, Release 41.1, Released on 6Jun2003, Formatted on 9Jun2003
SPTREMBL, Release 23.0, Released on 4Mar2003, Formatted on 7Mar2003

Total finds: 0
Total length: 305,079,309
Total sequences: 958,388
CPU time: 06:50.54

> 0 <
01:10 IntellGenetics
> 0 <

Quest - Quick User-directed Expression Search Tool
Release 5.4

-- Outline of search "seq6-1ss" --

Selected search type is key against sequence data banks or files.

Selected scope is Sequence.

Selected sequence key from "new.key":

seq6 (AA) ID seq6 AA preliminary pattern
1 followed by

2 c
r or k repeated 20 times

Selected data banks and files:

Data bank : Issued_AA , all entries

-- Output Parameters --

Format Options:

Nucleic acid code matching	Exact	File Options:	No
Find non-matching hits only	No	Indirect file	No
Report key used	Yes	Sequence or key file	No
Note position of hit	Yes	List of hits	No
Display full annotations	Yes	Hit display	Yes
Sequence context	50	Name and annotations	Yes

-- Run Parameters --

Run mode	Batch
Time to start comparison	now
Notify at end of run	No

No hits found.

-- Search Statistics --

Times:	CPU	Total Elapsed
	00:01:29.16	00:01:30.00
Number of sequences searched:		328807
Number of sequence hits:		0
Number of separate matches:		0
Number of sequence hits saved:		0

! FINDPATTERNS on pir:* allowing 0 mismatches

! 1 (R,K){20,20}

January 30, 2004 07:05 ..

NBRF, Release 76.1, Released on 12May2003, Formatted on 10Jun2003

Total finds: 17
Total length: 96,168,682
Total sequences: 283,308
CPU time: 03:26.44

1 T49173 ck: 4143 len: 517 ! hypothetical protein T20N10.250 - Arabidops

(R,K){20,20}

(K){20}

444: FERVG KKKKKKKKKKKKKKKKKKK KKKIR

(K){20}

445: ERVGR KKKKKKKKKKKKKKKKKKK KKIRL

(K){20}

446: RVGRK KKKKKKKKKKKKKKKKKKK KIRLN

(K){20}

447: VGRKK KKKKKKKKKKKKKKKKKKK IRLNF

1 T46395 ck: 7330 len: 380 ! hypothetical protein DKFZp434I1120.1 - huma

(R,K){20,20}

(K){20}

355: NLLIQ KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

356: LLLQK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

357: LLQKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

358: LQKKK KKKKKKKKKKKKKKKKKKK KKK

(K){20}

359: QKKKK KKKKKKKKKKKKKKKKKKK KK

(K){20}

360: KKKKK KKKKKKKKKKKKKKKKKKK K

(K){20}

361: KKKKK KKKKKKKKKKKKKKKKKKK

1 I52523 ck: 8048 len: 215 ! nucleoporin p62 homolog - rat (fragment)

(R,K){20,20}

(K){20}

35: CEPLE KKKKKKKKKKKKKKKKKKK KKTGD

(K){20}

36: EPLEK KKKKKKKKKKKKKKKKKKK KTGDN

(K){20}

37: FLEKK KKKKKKKKKKKKKKKKKKK TGDNA

1 S58321 ck: 1384 len: 126 ! probable membrane protein YOR309c - yeast

(R,K){20,20}

(K){20}

53: KRRTT RRRRRRRRRRRRRRRRRR KRSPR

(R,K){20}

54: KRRTT RRRRRRRRRRRRRRRRRR RSPRK

(R,K){20}

55: RRTTR RRRRRRRRRRRRRRRRRR SPRKR

Databases searched:


```
!!SEQUENCE_LIST 1.0
! FINDPATTERNS on plr:* allowing 0 mismatches
!      1 (R,K){20,20}      January 30, 2004 07:51 ..
```

```
PIR2:T49173      ck: 4143 len: 517 finds: 4      ! hypothetical protein T20N10.25
PIR2:T46395      ck: 7330 len: 380 finds: 7      ! hypothetical protein DKFZp434I
PIR2:I52523      ck: 8048 len: 215 finds: 3      ! nucleoporin p62 homolog - rat
PIR2:S58321      ck: 1384 len: 126 finds: 3      ! probable membrane protein YOR3
```

\\End of list

Databases searched:
NBRF, Release 76.1, Released on 12May2003, Formatted on 10Jun2003

```
Total finds:      17
Total length:    96,168,682
Total sequences: 283,308
CPU time:        03:53.65
```



```

!!AA SEQUENCE 1.0
P1:T49173 - hypothetical protein T20N10.250 - Arabidopsis thaliana
C1:Species: Arabidopsis thaliana (mouse-ear cress)
C1:Date: 02-Jun-2000 #sequence_revision 02-Jun-2000 #text_change 08-Dec-2000
C1:Accession: T49173
R1:D'Angelo, M.; Vezzi, A.; Modesto, D.; Pigazzi, M.; Valle, G.; Mewes, H.W.;
Rudd, S.; Lemcke, K.; Meyer, K.F.X.; Queclier, F.; Salanoubat, M.
submitted to the Protein Sequence Database, April 2000
A1:Reference number: Z25017
A1:Accession: T49173
A1:Status: preliminary
A1:Molecule type: DNA
A1:Residues: 1-517 <DNA>
A1:Cross-references: EMBL:AL353032; GSPDB:GN00061; ATSP:T20N10.250
A1:Experimental source: cultivar Columbia; BAC clone T20N10
C1:Genetics:
A1:Gene: ATSP:T20N10.250
A1:Map position: 3
A1:Introns: 312/3; 359/3; 444/3
C1:Superfamily: Arabidopsis thaliana hypothetical protein P17J16.30
T49173 Length: 517 January 30, 2004 07:58 Type: P Check: 4143 ..

1 MDLFSLEPNE LLYHLSFLS TKEALTSVL SKRWNLFAF VYLEPDDSV
51 FLHPERKRE KEGILQFMD FVDRVLDLHG DSLIKTFSLK CKTVGSDHV
101 DRWICNVLAR GVSDDLDFID FRDLYSLPHE VGVSRLLVVL RVGSSDLVW
151 WQFLCLPML KTLVLDSCWL CIGQFOILL ACPALBELDM TNRWQDSNV
201 TVSSSLKEL TIDHGCCSV VNLKSLFDA PSLVIFYCD SLAEDYPQVN
251 LKNLYBAQIN LLLTQAQIEQ VRALNEMLV ADDVFPGLGN AMKLITGLRN
301 VQQLYSPDT LEVSRCEG MPVFNNLKV LSIWSDNRGM QAMPVLLRNC
351 PHLETLIING LLYATDKCG DVCCISRDY KQSLTSCPV KKLQIYFRG
401 TIRELEMTGH FLKTPPCKE MDIYAHNSH TLFKDTTIE RVGKSKKKK
451 KKKKKKKKK KKKKKIRLN FKPVNKTQF LKRLADKLG IPQCLEPLDV
501 DSSLGELAIL AMDSRPS

!!AA SEQUENCE 1.0
P1:T46395 - hypothetical protein DKFZp43411120.1 - human (fragment)
C1:Species: Homo sapiens (man)
C1:Date: 04-Feb-2000 #sequence_revision 04-Feb-2000 #text_change 04-Feb-2000
C1:Accession: T46395
R1:Ottenwilder, B.; Obermaier, B.; Mewes, H.W.; Gassenhuber, J.; Wiemann, S.
submitted to the Protein Sequence Database, January 2000
A1:Reference number: Z23031
A1:Accession: T46395
A1:Status: preliminary
A1:Molecule type: mRNA
A1:Residues: 1-380 <AA>
A1:Cross-references: EMBL:AL137556
A1:Experimental source: adult testis; clone DKFZp43411120
C1:Genetics:
A1:Note: DKFZp43411120.1
T46395 Length: 380 January 30, 2004 07:58 Type: P Check: 7330 ..

1 WGSTDSKLNK RKAIVQLTTK TQPEATDDA FWDQFADTA TSVQDFALV
51 PAEIRAVRE ESPNLATLC YKAVEKLVQG AEGSGHSEKE KQIVNCSRL
101 LRRVLPYIRE DPMRGFFWS TVPAGRGGG EEDDEHARPL AESLLALAD
151 LIFCPDFTVQ SHRSTVDSA EDVHSLDSC YIWEAGVGR HSPQPNYIHD
201 NMRMELIKLL LTCFSEAMYL PPAEESGSTN PWQFCSTN NRHALPLFTS

```

```

!!AA SEQUENCE 1.0
P1:I52523 - nucleoporin p62 homolog - rat (fragment)
C1:Species: Rattus norvegicus (Norway rat)
C1:Date: 27-Feb-1997 #sequence_revision 27-Feb-1997 #text_change 05-Nov-1999
C1:Accession: I52523
R1:Wang, Z.Q.; Akmal, K.M.; Kim, K.H.
Biol. Reprod. 51, 1022-1030, 1994
A1>Title: An unusual nucleoporin-related messenger ribonucleic acid is present
in the germ cells of rat testis.
A1:Reference number: I52523; MUID:95151924; PMID:7849178
A1:Accession: I52523
A1:Status: preliminary; translated from GB/EMBL/DBJ
A1:Molecule type: mRNA
A1:Residues: 1-215 <RES>
A1:Cross-references: GB:575997; NID:g913245; PIDN:AA83384.1; PID:g913246
A1:Experimental source: testis
I52523 Length: 215 January 30, 2004 07:58 Type: P Check: 8048 ..

1 SGGKATSSCD EDCSSSLPF SLSGPVKQDC EFLKSKKKK KKKKKKKKK
51 KKKKKKTGDN AKSVSRQYSL KTKLEHEAE QAKVELDFIL SQKLELDL
101 SPLSEVKEQ SGTIYLQHAD EEREKTYKLA ENIDAQLKRM AQDKDIEH
151 LNMAGGPADT SDPLQOICKI LNMHMSLQW VQSSALLQR RVEASRYCE
201 SRKEQERSL RIAPD

!!AA SEQUENCE 1.0
P1:S58321 - probable membrane protein YOR309c - yeast (Saccharomyces cerevisiae)
M1:Alternate names: hypothetical protein O6105
C1:Species: Saccharomyces cerevisiae
C1:Date: 13-Jan-1996 #sequence_revision 01-Mar-1996 #text_change 19-Apr-2002
C1:Accession: S58321; S67215; S71989
R1:Pearson, B.M.; Hernando, Y.; Wolf, S.S.; Kalogeropoulos, A.; Schweizer, M.
submitted to the EMBL Data Library, August 1995
A1:Reference number: S58318
A1:Accession: S58321
A1:Molecule type: DNA
A1:Residues: 1-126 <PEA>
A1:Cross-references: EMBL:X90565; NID:g940836; PID:g940840
R1:Pearson, B.M.; Hernando, Y.; Kalogeropoulos, A.; Schweizer, M.
submitted to the Protein Sequence Database, July 1996
A1:Reference number: S67213
A1:Accession: S67215
A1:Molecule type: DNA
A1:Residues: 1-126 <PEW>
A1:Cross-references: EMBL:Z75217; NID:g1420680; PID:e252431; PID:g1420681;
MIPS:YOR309C
A1:Experimental source: strain S288C
R1:Pearson, B.M.; Hernando, Y.; Payne, J.; Wolf, S.S.; Kalogeropoulos, A.;
Schweizer, M.
Yeast 12, 1021-1031, 1996
A1>Title: Sequencing of a 35.71 kb DNA segment on the right arm of yeast
chromosome XV reveals regions of similarity to chromosomes I and XIII.
A1:Reference number: S71989; MUID:97051589; PMID:8896266
A1:Accession: S71989
A1:Status: nucleic acid sequence not shown; translation not shown
A1:Molecule type: DNA
A1:Residues: 1-126 <PEF>
A1:Cross-references: EMBL:X90565; NID:g940836; PIDN:CA62164.1; PID:g940840
A1:Note: the nucleotide sequence was submitted to the EMBL Data Library, August
1995
C1:Genetics:
A1:Cross-references: SGD:S0005836

```

A:Map position: 15R
 C:Keywords: transmembrane protein
 F:3-19/Domain: transmembrane #status predicted <TM1>
 F:107-123/Domain: transmembrane #status predicted <TM2>
 SS8321 Length: 126 January 30, 2004 07:58 Type: P Check: 1384 ..
 1 MQMLIPQRL LILNPLMMK RKKRKKRKR RERETWMLP RILKKLRKR
 51 RTRRKKRKR KRRRKKRKR RKRSPRKR KRNKDAFYI LIIDPSRSL
 101 LRGFRKFSII IQCLTYFSFH ILFHNL

FINDPATTERNS on swp:* allowing 0 mismatches
1 (R,K){20,20} January 30, 2004 07:00 ..

1 Q12444 ck: 1384 len: 126 1 Q12444 saccharomyces cerevisiae (baker's ye
(R,K){20,20}
(R,K){20}
53: RKRRT RRRRRRRRRRRRRRRRRR KRSPR
(R,K){20}
54: KRRTT RRRRRRRRRRRRRRRRRR RSPRK
(R,K){20}
55: RRTTR RRRRRRRRRRRRRRRRRR SPRKR

1 Q9P529 ck: 291 len: 128 1 Q9P529 neurospora crassa. hypothetical 15.2
(R,K){20,20}
(K){20}

71: KRNQ KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
72: RKNQ KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
73: KNQK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
74: NQKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
75: QKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
76: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
77: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
78: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
79: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
80: KKKK KKKKKKKKKKKKKKKKKKK KKEQ
(K){20}
81: KKKK KKKKKKKKKKKKKKKKKKK KKEQ
(K){20}
82: KKKK KKKKKKKKKKKKKKKKKKK KQES
(K){20}
83: KKKK KKKKKKKKKKKKKKKKKKK EDES

1 Q9NT34 ck: 7330 len: 380 1 Q9nt34 homo sapiens (human). hypothetical 1
(R,K){20,20}
(K){20}

355: NLLQ KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
356: LLLQ KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
357: LLQK KKKKKKKKKKKKKKKKKKK KKKK

358: LQKK KKKKKKKKKKKKKKKKKKK KK
(K){20}
359: QKKK KKKKKKKKKKKKKKKKKKK KK
(K){20}
360: KKKK KKKKKKKKKKKKKKKKKKK K
(K){20}
361: KKKK KKKKKKKKKKKKKKKKKKK

1 Q9H6Q7 ck: 3351 len: 720 1 Q9h6q7 homo sapiens (human). hypothetical 1
(R,K){20,20}
(K){20}

692: IVSIS KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
693: VSISK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
694: SISK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
695: ISKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
696: SKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
697: KKKK KKKKKKKKKKKKKKKKKKK KKK
(K){20}
698: KKKK KKKKKKKKKKKKKKKKKKK KK
(K){20}
699: KKKK KKKKKKKKKKKKKKKKKKK KK
(K){20}
700: KKKK KKKKKKKKKKKKKKKKKKK K
(K){20}
701: KKKK KKKKKKKKKKKKKKKKKKK

1 Q8N6F0 ck: 9898 len: 55 1 Q8n6f0 homo sapiens (human). similar to loc
(R,K){20,20}
(R,K){20}

14: RRGK KKKRRRRRRRRRRRRRRRRR KKKK
(R,K){20}
15: RRGK KKKRRRRRRRRRRRRRRRRR KKKK
(R,K){20}
16: GRGK KKKRRRRRRRRRRRRRRRRR KKKK
(R,K){20}
17: RGGK KKKRRRRRRRRRRRRRRRRR KKKK
(R,K){20}
18: GKKK RRRRRRRRRRRRRRRRRRR KKKK
(R,K){20}
19: KKKK KKKKKKKKKKKKKKKKKKK KKKR
(R,K){20}
20: KKKK RRRRRRRRRRRRRRRRRRR KRRR
(K){20}
21: KRRR KKKKKKKKKKKKKKKKKKK KRRR

22: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK RRRRR
23: RRRKK (R,K) {20} KKKKKKKKKKKKKKKKKKK RRRRR
24: KKKKK (R,K) {20} KKKKKKKKKKKKKKKKKKK RRRRG
25: RKKKK (R,K) {20} KKKKKKKKKKKKKKKKKKK RRRGR
26: KKKKK (R,K) {20} KKKKKKKKKKKKKKKKKKK RRRGR
27: KKKKK (R,K) {20} KKKKKKKKKKKKKKKKKKK RRRRR
28: KKKKK (R,K) {20} KKKKKKKKKKKKKKKKKKK RRRRR

1 Q9HC48 ck: 7602 len: 667 1 Q9hc48 homo sapiens (human). ctcl tumor ant
(R,K) {20,20}
648: GDXTD RKKKKKKKKKKKKKKKKKKKK

1 Q9H5V6 ck: 379 len: 168 1 Q9h5v6 homo sapiens (human). hypothetical p
(R,K) {20,20}
140: VREME KKKKKKKKKKKKKKKKKKKKK KKKKK
141: REMEK KKKKKKKKKKKKKKKKKKKKK KKKKK
142: EWEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
143: WEKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
144: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
145: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
146: KKKKK KKKKKKKKKKKKKKKKKKKKK KKK
147: KKKKK KKKKKKKKKKKKKKKKKKKKK KK
148: KKKKK KKKKKKKKKKKKKKKKKKKKK K
149: KKKKK KKKKKKKKKKKKKKKKKKKKK

1 Q8SWR7 ck: 2372 len: 515 1 Q8swr7 drosophila melanogaster (fruit fly).
(R,K) {20,20}
493: FTDYI KKKKKKKKKKKKKKKKKKKKK KKK
494: TDYIK KKKKKKKKKKKKKKKKKKKKK KK

495: DYIKK KKKKKKKKKKKKKKKKKKKKK K
496: YIKKK KKKKKKKKKKKKKKKKKKKKK
Q8T2U7 ck: 8768 len: 791 1 Q8t2u7 dictyostelium discoideum (slime mold)
(R,K) {20,20}
769: EIEKE KKKKKKKKKKKKKKKKKKKKK KEI
770: IEKEK KKKKKKKKKKKKKKKKKKKKK EI

1 Q8I247 ck: 5951 len: 206 1 Q8i247 plasmodium falciparum (isolate 3d7).
(R,K) {20,20}
185: KKIPL KKKKKKKKKKKKKKKKKKKKK KT
186: KIFLK KKKKKKKKKKKKKKKKKKKKK T

1 Q95LV6 ck: 7515 len: 531 1 Q95lv6 macaca fascicularis (crab eating mac
(R,K) {20,20}
502: YKNS KKKKKKKKKKKKKKKKKKKKK KKKKK
503: KNSKK KKKKKKKKKKKKKKKKKKKKK KKKKK
504: GNSKK KKKKKKKKKKKKKKKKKKKKK KKKKK
505: NSKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
506: SKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
507: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
508: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
509: KKKKK KKKKKKKKKKKKKKKKKKKKK KKK
510: KKKKK KKKKKKKKKKKKKKKKKKKKK KK
511: KKKKK KKKKKKKKKKKKKKKKKKKKK K
512: KKKKK KKKKKKKKKKKKKKKKKKKKK

1 Q9LXR2 ck: 4143 len: 517 1 Q9lxr2 arabidopsis thaliana (mouse-ear cress)
(R,K) {20,20}
444: FERVG KKKKKKKKKKKKKKKKKKKKK KKKIR
445: ERVGK KKKKKKKKKKKKKKKKKKKKK KKIRL

446: RVGKK (K){20} KKKKKKKKKKKKKKKKK KIRLN
447: VGGKK (K){20} KKKKKKKKKKKKKKKKK IRLNF

1 O8S7D3 ck: 6479 len: 80 i O8s7d3 oryza sativa (rice). hypothetical 9.

48: VIHLD (R,K){20,20} KKKKKKKKKKKKKKKKK KKKKK
(K){20}
49: IHLDK (K){20} KKKKKKKKKKKKKKKKK KKKKK
50: HLDKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
51: LDGKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
52: DKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
53: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
54: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKL
55: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKLGL
56: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKLGE
57: KKKKK (K){20} KKKKKKKKKKKKKKKKK LIGE

1 Q8LP6 ck: 3239 len: 113 i Q8lp6 oryza sativa (japonica cultivar-grou

10: SLEHT (R,K){20,20} KKKKKKKKKKKKKKKKK KKKKK
(K){20}
11: LEHIK (K){20} KKKKKKKKKKKKKKKKK KKKKK
12: EHIKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
13: HIKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
14: IKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
15: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
16: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
17: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
18: KKKKK (R,K){20} KKKKKKKKKKKKKKKKK KKKKK

Q9IG29 ck: 6094 len: 260 i Q9lg29 arabidopsis thaliana (mouse-ear cres

1

6: MDRCI (R,K){20,20} KKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
7: DRCIR (K){20} KKKKKKKKKKKKKKKKK KKKKK
8: RCIRK (K){20} KKKKKKKKKKKKKKKKK KKKKK
9: CIRKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
10: IRKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
11: RKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
12: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
13: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
14: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
15: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
16: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
17: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
18: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
19: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
20: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
21: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
22: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
23: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
24: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
25: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
26: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
27: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
28: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK
29: KKKKK (K){20} KKKKKKKKKKKKKKKKK KKKKK

127: KKKK (K) {20} KKKK
 128: KKKK (K) {20} KKKK
 129: KKKK (K) {20} KKKK
 130: KKKK (K) {20} KKKK
 131: KKKK (K) {20} KKKK
 132: KKKK (K) {20} KKKK
 133: KKKK (K) {20} KKKK
 134: KKKK (K) {20} KKKK
 135: KKKK (K) {20} KKKK
 136: KKKK (K) {20} KKKK
 137: KKKK (K) {20} KKKK
 138: KKKK (K) {20} KKKK
 139: KKKK (K) {20} KKKK
 140: KKKK (K) {20} KKKK
 141: KKKK (K) {20} KKKK
 142: KKKK (K) {20} KKKK
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 145: KKKK (K) {20} KKKK
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 147: KKKK (K) {20} KKKK
 148: KKKK (K) {20} KKKK
 149: KKKK (K) {20} KKKK
 150: KKKK (K) {20} KKKK

151: KKKK KKKK KKKK KKKK KKKK
 152: KKKK (K) {20} KKKK
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222: KKKK (K) {20}
KKKK
223: KKKK (K) {20}
KKKK

224: KKKKK KKKKKKKKKKKKKKKKKKK KKKKR
(K) {20}
225: KKKKK KKKKKKKKKKKKKKKKKKK KKKRH
(K) {20}
226: KKKKK KKKKKKKKKKKKKKKKKKK KKRHH
(K) {20}
227: KKKKK KKKKKKKKKKKKKKKKKKK KKRHH
(K) {20}

Q64075 ck: 8048 len: 215 ! Q64075 rattus sp. nucleoporin p62 homolog R
(R,K) {20,20}
(K) {20}
35: CEFLK KKKKKKKKKKKKKKKKKKK KKTGD
(K) {20}
36: EFLEK KKKKKKKKKKKKKKKKKKK KTGDN
(K) {20}
37: FLEKK KKKKKKKKKKKKKKKKKKK TGDNA
(K) {20}

Q9D5G1 ck: 9398 len: 169 ! Q9D5G1 mus musculus (mouse). adult male tes
(R,K) {20,20}
(R) {20}
117: VOLRG RRRRRRRRRRRRRRRRRR RRRKK
(R) {20}
118: QLRGR RRRRRRRRRRRRRRRR RKKKE
(R) {20}
119: LRGRR RRRRRRRRRRRRRRRR RKKKE
(R) {20}
120: RGRRR RRRRRRRRRRRRRRRR RKEEE
(R,K) {20}
121: GRRRR RRRRRRRRRRRRRRRR KEEEE
(R,K) {20}
122: RRRRR RRRRRRRRRRRRRRRR EEEEE
(R,K) {20}

Q35807 ck: 7510 len: 129 ! Q35807 rattus norvegicus (rat). microvascul
(R,K) {20,20}
(K) {20}
85: VLLAS KKKKKKKKKKKKKKKKK KKKKE
(K) {20}
86: LLASK KKKKKKKKKKKKKKKKK KKKKE
(K) {20}

Q8BXG9 ck: 5434 len: 115 ! Q8BXG9 mus musculus (mouse). hypothetical a
(R,K) {20,20}
(R,K) {20}
40: IIRRR RRRRRRRRRRRRRRRR KETGS
(R,K) {20}
41: IIRRR RRRRRRRRRRRRRRRR ETGSH
(R,K) {20}

Q8BHV2 ck: 8958 len: 154 ! Q8bhv2 mus musculus (mouse). weakly similar
(R,K) {20,20}
(R) {20}

42: RGEE RRRRRRRRRRRRRRRR RRRR

```

43: RGEER RRRRRRRRRRRRRRRRRR RRKR
      (R) {20}

44: GEER RRRRRRRRRRRRRRRRRR RRKR
      (R) {20}

45: EERR RRRRRRRRRRRRRRRRRR RRKR
      (R) {20}

46: ERRR RRRRRRRRRRRRRRRRRR RRKR
      (R) {20}

47: RRRR RRRRRRRRRRRRRRRRRR RRKR
      (R) {20}

48: RRRR RRRRRRRRRRRRRRRRRR RRKR
      (R) {20}

49: RRRR RRRRRRRRRRRRRRRRRR RRKE
      (R) {20}

50: RRRR RRRRRRRRRRRRRRRRRR RKER
      (R) {20}

51: RRRR RRRRRRRRRRRRRRRRRR KRER
      (R) {20}

52: RRRR RRRRRRRRRRRRRRRRRR RRRR
      (R,K) {20}

53: RRRR RRRRRRRRRRRRRRRRRR ERRE
      (R,K) {20}

```

```

Databases searched:
  SWISS-PROT, Release 41.1, Released on 6Jun2003, Formatted on 9Jun2003
  SPTKMBE, Release 23.0, Released on 4Mar2003, Formatted on 7Mar2003

Total finds:      348
Total length:    305,079,309
Total sequences: 958,388
CPU time:        11:25.82

```

```
!!SEQUENCE LIST 1.0
! FINDPATTERNS on swp:* allowing 0 mismatches
!      1 (R,K){20,20}
```

January 30, 2004 07:59 ..

```
SP_FUN:Q12444      ck: 1384 len: 126 finds: 3      ! Q12444 saccharomyces cerevisiae
SP_FUN:Q9P529      ck: 291 len: 128 finds: 13      ! Q9P529 neurospora crassa. hyc
SP_HUM:Q9NT34      ck: 7330 len: 380 finds: 7       ! Q9NT34 homo sapiens (human). h
SP_HUM:Q9H6Q7      ck: 3351 len: 720 finds: 10      ! Q9H6Q7 homo sapiens (human). h
SP_HUM:Q8N6F0      ck: 9898 len: 55 finds: 15      ! Q8N6F0 homo sapiens (human). s
SP_HUM:Q9HC48      ck: 7602 len: 667 finds: 1       ! Q9HC48 homo sapiens (human). d
SP_HUM:Q9H5V6      ck: 379 len: 168 finds: 10      ! Q9H5V6 homo sapiens (human). h
SP_IN:Q8SWR7       ck: 2372 len: 515 finds: 4       ! Q8SWR7 drosophila melanogaster
SP_IN:Q8T2U7       ck: 8768 len: 791 finds: 2       ! Q8T2U7 dictyostelium discoideu
SP_IN:Q8I247       ck: 5951 len: 206 finds: 2       ! Q8I247 plasmodium falciparum
SP_OM:Q95LV6       ck: 7515 len: 531 finds: 11      ! Q95LV6 macaca fascicularis (cr
SP_PL:Q9LXR2       ck: 4143 len: 517 finds: 4       ! Q9LXR2 arabidopsis thaliana (m
SP_PL:Q8S7D3       ck: 6479 len: 80 finds: 10      ! Q8S7D3 oryza sativa (rice). hy
SP_PL:Q8LQP6       ck: 3239 len: 113 finds: 9       ! Q8LQP6 oryza sativa (japonica
SP_PL:Q9IGZ9       ck: 6094 len: 260 finds: 222     ! Q9IGZ9 arabidopsis thaliana (m
SP_RO:Q64075       ck: 8048 len: 215 finds: 3       ! Q64075 rattus sp. nucleoporin
SP_RO:Q9D5G1       ck: 9388 len: 169 finds: 6       ! Q9D5G1 mus musculus (mouse). a
SP_RO:Q35807       ck: 7510 len: 129 finds: 2       ! Q35807 rattus norvegicus (rat)
SP_RO:Q8BXG9       ck: 5434 len: 115 finds: 2       ! Q8BXG9 mus musculus (mouse). h
SP_RO:Q8BHV2       ck: 8958 len: 154 finds: 12      ! Q8BHV2 mus musculus (mouse). w
```

\\End of list

Databases searched:

SWISS-PROT, Release 41.1, Released on 6jun2003, Formatted on 9jun2003
 SPTREMBL, Release 23.0, Released on 4mar2003, Formatted on 7mar2003

Total finds: 348
 Total length: 305,079,309
 Total sequences: 958,388
 CPU time: 12:54.92


```

!!AA_SEQUENCE 1.0
ID_012444 PRELIMINARY; PRT; 126 AA.
AC 012444;
DT 01-NOV-1996 (TReMBLrel. 01, Created)
DT 01-NOV-1996 (TReMBLrel. 01, Last sequence update)
DT 01-OCT-2002 (TReMBLrel. 22, Last annotation update)
DE ORF YOR309C.
GN YOR309C.
OS Saccharomyces cerevisiae (Baker's Yeast).
OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
OC Saccharomycetales; Saccharomycetaceae; Saccharomyces.
OX NCBI_TaxID=4932;
RN [1]
RP SEQUENCE FROM N.A.
RA Pearson B.M., Hernando Y., Kalogeropoulos A., Schweizer M.;
RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RA MIPS;
RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=F1673;
RA Pearson B.M., Hernando Y., Wolf S.S., Kalogeropoulos A., Schweizer M.;
RL Submitted (AUG-1995) to the EMBL/GenBank/DBJ databases.
DR EMBL; 275217; CAA99629.1; -.
DR EMBL; X90565; CAA62164.1; -.
DR SGP; S0005836; YOR309C.
SQ SEQUENCE 126 AA; 16294 MW; 46E1F4C66480208 CRC64;

Q12444 Length: 126 January 30, 2004 08:18 Type: P Check: 1384 ..

1 MGLMPLPRL LILNPLMMK RKKKKKKKK RERETMMKIP RILKLRKKR
51 RTRRRRRRR RRRRRRRRR RRRSPRRR KRRNDAYFI LIISPSRSL
101 LFGFRKFSII IQCLTFYSFH ILFHNH

!!AA_SEQUENCE 1.0
ID_09529 PRELIMINARY; PRT; 128 AA.
AC 09529;
DT 01-OCT-2000 (TReMBLrel. 15, Created)
DT 01-DEC-2001 (TReMBLrel. 19, Last sequence update)
DT 01-OCT-2002 (TReMBLrel. 22, Last annotation update)
DE Hypothetical 15.2 kDa protein.
GN B24H17.160.
OS Neurospora crassa.
OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Sordariomycetes;
OC Sordariales; Sordariaceae; Neurospora.
OX NCBI_TaxID=5141;
RN [1]
RP SEQUENCE FROM N.A.
RA Schulte U., Aign V., Hohenel J., Brandt P., Fartmann B., Holland R.,
RA Nykatura G., Mewes H.W., Manhaupt G.;
RL Submitted (MAY-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RA German Neurospora genome project;
RL Submitted (OCT-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AL356815; CAB92638.2; -.
KW Hypothetical protein.
SQ SEQUENCE 128 AA; 15157 MW; 8C7C65C3DFB70765 CRC64;

Q9529 Length: 128 January 30, 2004 08:18 Type: P Check: 291 ..

1 MAISIGLHH KNINRRAPGH SVYSKSSYD PQQYDATOHY LPSQGFKAIP
51 DLITGGKQGC LSTHDKRNG KKKKKKKKK KKKKKKKKK KKKKKKKKK
101 KKEQSRITF QOHFOADGIC PTPWHTTR

!!AA_SEQUENCE 1.0
ID_09NT34 PRELIMINARY; PRT; 380 AA.

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AC 09NT34;
DT 01-OCT-2000 (TReMBLrel. 15, Created)
DT 01-OCT-2000 (TReMBLrel. 15, Last sequence update)
DT 01-OCT-2002 (TReMBLrel. 22, Last annotation update)
DE Hypothetical protein (Fragment).
GN DKF2P4341120.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Testis;
RA Oltewelder B., Obermaier B., Mewes H.W., Gaassenhuber J., Wiemann S.;
RL Submitted (JAN-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AL137556; CAB70810.1; -.
DR Genew; HGNC:15736; C17orf28.
KW Hypothetical protein.
FT NON TER 380
SQ SEQUENCE 380 AA; 42689 MW; 67F50DD101346AFB CRC64;

Q9NT34 Length: 380 January 30, 2004 08:18 Type: P Check: 7330 ..

1 MGSTDSKLNK RKAVIQUTTK TQPEATDPA FWDQFWADTA TSVQDVFAIV
51 PAAEIRAVRE ESPENLALIC YKAVELYQG AESGCHSEK KQIVLNCRL
101 LTRVLPYTFE DPMWRGFFWS TYFGAGRGG EEDDHARPL AESLILAIAD
151 LIFCPDFTVQ SHRRSTVDSA EDVHSLDSCB YIWEAGVGA HSPQPNYIHD
201 MNRMELKLIL LTCFSEAMYL PRAPESGSTN PWVQFCSTG NRHMLPLFTS
251 LINTVCAYDP VGIGIPYNNH LFSVREPLV EEAQVLIYT LDHDSASSAS
301 PTVDGTTTGT AMDADDPGP ENLFVNYLSR IHREDFQFI LKGIARLLSN
351 LLLQKKKKKK KKKKKKKKK KKKKKKKKK

!!AA_SEQUENCE 1.0
ID_09H6Q7 PRELIMINARY; PRT; 720 AA.
AC 09H6Q7;
DT 01-MAR-2001 (TReMBLrel. 16, Created)
DT 01-MAR-2001 (TReMBLrel. 16, Last sequence update)
DT 01-OCT-2002 (TReMBLrel. 22, Last annotation update)
DE Hypothetical protein FLJ21979 (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Kawabata A., Hiki J. T., Kobatake N., Inagaki H., Ikema Y., Okamoto S.,
RA Oktani R., Ota T., Suzuki Y., Oiyashi M., Nishi T., Shibahara T.,
RA Tanaka T., Nakamura Y., Isegaki T., Sugano S.;
RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AK025632; BAB15196.1; -.
KW Hypothetical protein.
FT NON TER 720
SQ SEQUENCE 720 AA; 84029 MW; A86586FEAA953D0B CRC64;

Q9H6Q7 Length: 720 January 30, 2004 08:18 Type: P Check: 3351 ..

1 MLTEQVEQYT KEMKNTCII EDLKNELQRN KGASTLSQQT HMKIQSTLDI
51 LKERTKEAR TAELEADAR EKDKELVEAL KKLKDYEGV YGLDAVVEI
101 KNCKNQIKIR DREIEITKE INKLELKISD FLDEBAARE RVGLEPKTMI
151 DLTEFRNSKH LKQOQYRAEN QILKIEISL EERLDLKKK IRQWAQERK
201 RSATSGLTTE DLNLTENISQ GDRISERKLD LLSLKNMSEA QSKNEFLSRE

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251 LIEKERDLER SRTYIAFQK KLKELVEENK QLEEGKEIT QAIKEMQKP
301 DVKGGETSLI IPSLERLVNA IESNNAEGIF DASLHLKAQV DQLTRENEEL
351 ROELRESRKE AINYSQOLAK ANLIKIDHLEK ETSILRQSEB SNVVEKGIDL
401 PGDIAPSSAS IINSQNEYLI HLLQELEENKE KKLKNELEDSL EDYNNKFAVI
451 RHQOSILYKE YLSEKETWKT ESKITKEER KLEDOYQODA IKVKEYNNLL
501 NALQWDSDEM KKIILAENSRK ITVLQVNEKS LIRQYTTLVE LERQURKENE
551 KQKNELLSME AEVECKIGCL QRFKEMAFK IALQKVON SVLSSELELA
601 NKQYNEITAK YRDILQKDNM LVQRTSNLEH LECENISLKE QVESINKLE
651 ITREKLHTIE QAMEQETKLG NESSMDRAKK SITNSDIVSI SKKKKKKKKK
701 KKKKKKKKKK KKKKKKKKKK

!!AA_SEQUENCE 1.0
ID_Q8N6F0 PRELIMINARY; PRT; 55 AA.
AC_Q8N6F0;
DT 01-OCT-2002 (TREMBLrel. 22, Created)
DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Similar to LOC201361.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
OX NCBI_TaxId=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Lung;
RA Strausberg R.;
RL Submitted (May-2002) to the EMBL/Genbank/DBJ databases.
DR EMBL; BC030525; AAH30525.1; -
SQ SEQUENCE 55 AA; 7251 MW; 0906032B284006BA CRC64;

Q8N6F0 Length: 55 January 30, 2004 08:18 Type: P Check: 9898 ..

1 MFLASQRERR GRGKKKKRKR KKKKKKKKKK KKKKKKKKKK KRRRRRGRGR
51 RRMQO

!!AA_SEQUENCE 1.0
ID_Q9HC48 PRELIMINARY; PRT; 667 AA.
AC_Q9HC48;
DT 01-MAR-2001 (TREMBLrel. 16, Created)
DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE CTCL tumor antigen se2-5 (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
OX NCBI_TaxId=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Testis;
RX MEDLINE=21143360; PubMed=1149944;
RA Eichmuller S.; Usener D.; Dummer R.; Stein A.; Thiel D.;
RA Schendendorf D.;
RT "Serological detection of cutaneous T-cell lymphoma-associated
antigens."
RL Proc. Natl. Acad. Sci. U.S.A. 98:629-634(2001).
CC -1- SIMILARITY: CONTAINS 2 PDZ/DHR DOMAINS.
DR EMBL; AF177228; AAG33676.1; -
DR HSSP; Q12923; 3PDZ.
DR InterPro; IPR001478; PDZ.
DR Pfam; PF00595; PDZ; 2.
DR SMART; SMO0228; PDZ; 2.
DR PROSITE; PSS0106; PDZ; 2.

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FT NON TER 1 1
FT NON TER 667 667
SQ SEQUENCE 667 AA; 73499 MW; C653EC16802BAE02 CRC64;

Q9HC48 Length: 667 January 30, 2004 08:18 Type: P Check: 7602 ..

1 EHENLFREND CIVRINDDL RNRFEQAOH MERQAMRTPI IMFHVPAAN
51 KEQYEQLSQS EKNNYSSRF SPDQYIDNR SVNSAGLHTV QVARRLNHPP
101 EOIDSHSRLP HSAHPGKPP SAPASAPQNV FSTTVSSGVN TKIKGKRLNI
151 QLKGTGEGLE FSITSRDVTI GGSAPIYVKN ILPRGAIQD GRLXAGDRLI
201 EVNGVDLVGK SQEEVSLR STMEGTVSL LVFRQEDAFH PRELNAPSQ
251 MQIPREYAE DEDIVLTPDG TREFLTPEVP LNDSSAGLG VSVGNRSKE
301 NHADLGIYK SIINGASK DGLRYNDQL IAVNGESLLG KTNQDAMETL
351 RRMSTEGNK RGMQLIVAR RISKNEKLS PGSPGPELP IETALDRER
401 RISHSLYSGI EGLDESPSRN AALSRIWGS GKYLSPVTN MPQDDTYIIE
451 DRLFLVLPFH LSDQSSSSH DDVGFYTADA GTWAKAASD SADCSLSPDV
501 DPLVAFQREG FGRLADETK LNTVDDQKAG SPSRDVGPGL GLKXSSLSLS
551 LOTVAEYTL NGDIPFHRPR PRIIRGCGN ESFRAIDKS YDKAVDDDD
601 EGMETLEBDT EESSRSGRES VSTASDOPSH SLERQWNGNQ EKGDKTRKK
651 KKKKKKKKKK KKKKKKKK

!!AA_SEQUENCE 1.0
ID_Q9H5V6 PRELIMINARY; PRT; 168 AA.
AC_Q9H5V6;
DT 01-MAR-2001 (TREMBLrel. 16, Created)
DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
DT 01-OCT-2002 (TREMBLrel. 22, Last annotation update)
DE Hypothetical protein FLJ22976 (Fragment).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
OX NCBI_TaxId=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Watanabe K.; Kumagai A.; Itakura S.; Yamazaki M.; Tashiro H.; Ota T.;
RA Suzuki Y.; Oiyashi M.; Nishi T.; Shibahara T.; Tanaka T.;
RA Nakamura Y.; Isegai T.; Sugano S.;
RT "NEBO human cDNA sequencing project."
RL Submitted (AUG-2000) to the EMBL/Genbank/DBJ databases.
DR EMBL; AK026629; BAB15513.1; -
KW Hypothetical protein.
FT NON TER 168 168
SQ SEQUENCE 168 AA; 19549 MW; A19BDB195F8A1A90 CRC64;

Q9H5V6 Length: 168 January 30, 2004 08:18 Type: P Check: 379 ..

1 MNGNGRSGIQ QKGKNDGVA ATPAASAC QYRCIECNOE AKELYRDYNH
51 GVLKTTICKS CQKPVDKYIE YDPVILINA ILCKAQAYRH ILFTVQINIH
101 GKLCTFLCLC EAYLRWQLO DSNQNTAPPD LIRYRENEK KKKKKKKKKK
151 KKKKKKKKKK KKKKKKKK

!!AA_SEQUENCE 1.0
ID_Q8SWR7 PRELIMINARY; PRT; 515 AA.
AC_Q8SWR7;
DT 01-JUN-2002 (TREMBLrel. 21, Created)
DT 01-JUN-2002 (TREMBLrel. 21, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)

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DE GH22607p (Fragment).
GN CG7180.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;
OC Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
OX NCBI_TaxID=7227;
RN
RP SEQUENCE FROM N.A.
RC STRAIN=Berkely.
RA Stapleton M., Brokstein P., Hong L., Agbayani A., Carlson J.,
RA Change M., Chavez C., Dorsett V., Drensek D., Fartin D., Frise E.,
RA George R., Gonzalez M., Guarin H., Krommiller B., Li P., Liao G.,
RA Miranda A., Mungall C.J., Nuno C., Pacleb J., Paragas V., Park S.,
RA Patel S., Phouanavong S., Wan K., Yu C., Lewis S.E., Rubin G.M.,
RA Celinker S.;
RL Submitted (Apr-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY095518; AAM1251.1; -.
DR FlyBase; FBgn003673; CG7180.
DR InterPro; IPR000387; TYR_phosphatase.
DR InterPro; IPR000242; TYR_PP.
DR Pfam; PF00102; Y_phosphatase; 1.
DR SMART; SM00194; PTPc; 1.
DR PROSITE; PS00383; TYR_PHOSPHATASE_1; 1.
DR PROSITE; PS50056; TYR_PHOSPHATASE_2; 1.
DR PROSITE; PS50055; TYR_PHOSPHATASE_PTP; 1.
KM Hydroxylase.
KW
SQ
FT NON TER 515 515
FT SEQUENCE 515 AA; 59080 MW; B2825B7EA961958 CRC64;
Q8SWR7 Length: 515 January 30, 2004 08:18 Type: P Check: 2372 ..

1 MWTQQLVGC PEALNEEKS PAASAAGA GADMTAATGG GSSGAGGK
51 GGRSRSSAR YDDVEKQQR SRATVSPNT IKLMSLNG LSPERIKLEA
101 RDENSLSKT IPNGPIDRH FLKLDLRRK PVLTKLEFQ TAAKVESNTC
151 RLAKKNLE KNQPKCIPY DYNRVLEKV GGLQSDSYN ASYVDSLAKP
201 NAYITQGV EETVOAYWRM WQENISAIV MLTKTFPAK VMCHOYMPN
251 MEVHQYGD FINIVREQL ANFHITPRL YKNEKEVLT DERLILQPHY
301 TEWYHSCPF SNALLEFRR VRLVGNIIK DEDDMGPII VHCSGGGERS
351 GYVMSIDANL ELAEEBCFN VFGLKLRQ SRKGLVENVE QYKITYTLE
401 EHIGKTFV PVSEISDRK AKARRNSGK MNEYQAEDQ ICKQTPFTI
451 GDCAGHRAD NREKNRDLV VPDPNRPYL TSGQNAFTD YIKKKKKKKK
501 KKKKKKKKK KKKKK
11AA SEQUENCE 1.0
ID O8T2U7 PRELIMINARY; PRT; 791 AA.
AC O8T2U7;
DT 01-JUN-2002 (Tremblrel. 21, Created)
DT 01-JUN-2002 (Tremblrel. 21, Last sequence update)
DT 01-MAR-2003 (Tremblrel. 23, Last annotation update)
DE Hypothetical 92.4 kDa protein.
OS Dictyostelium discoideum (Slime mold).
OC Eukaryota; Mycetozoa; Dictyostelida; Dictyostelium.
OX NCBI_TaxID=44689;
RN
RP SEQUENCE FROM N.A.
RC STRAIN=X4;
RA Gloeckner G., Eichinger L., Szafranski K., Pachebat J., Dear P.,
RA Lehmann R., Baumgart C., Parra G., April J.F., Guigo R., Kumpf K.,
RA Tunga B., Cox E., Quail M.A., Platzer M., Rosenchal A., Noegel A.A.;
RT "Sequence and Analysis of Chromosome 2 of Dictyostelium";
RL Submitted (Mar-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC115574; AAL92183.1; -.

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DR InterPro; IPR005033; YEATS.
DR InterPro; IPR007087; ZnF_C2H2.
DR Pfam; PF03366; YEATS; 1.
DR SMART; SM00355; ZnF_C2H2; 1.
KM Hypothetical protein.
SQ
FT NON TER 791 AA; 92375 MW; D66C6BDEC92352C CRC64;
Q8T2U7 Length: 791 January 30, 2004 08:18 Type: P Check: 8768 ..

1 MEKETETLL DTIDEKENS TTTTQNTNH NNNNTNTNN NNNNNNNN
51 NNNNNNNNN NNNNNNNNN NNPSTLSAS LSKLRKLI NNEGDRELK
101 KRLIMIRE PPAELKKEQ ELIEIDRLN TYKEMLSLN KQNSTKST
151 YHIFTNNNN NNNNNNNNN NNNNNNNNN NNNNNNNNN NNNNNNNNN
201 NKKPRIFHK LQNGEPVLY CKYCRSDPV SGLGFLNHR IKHGFYSTL
251 DEARQIGVP VADSEIPKD PSREGIVFP KGAVSPSIA FERNSEEDNG
301 TFDNNNNNN NNNNNNNNN NNNNNNTGN DTDKKNNG NDADVDIDL
351 NVNSYNNK ETEESSGS RPYVKKIIV GNTSTQHPD YRGHDSRHK
401 WTVYRGPN EADISFVK IPEYLHSPA PDKVEYVER PFNLTRGNG
451 EPPVRIPLF HDKKNRPDI IHLKLIQL IQYVPVVG ETTETIDLR
501 LFFDKRQQL KLDQLNNNN NNNSENNN NNNNNNNNN NNINNNNN
551 INNNNDNN SSNTSPTSN YLNEPKIVN NDKVSENS NEDSQKXK
601 EKEKEKEK EKEKEKEK EKEKDKK EKEKDKK VKENDKKE
651 LNNRPKDR DRDEREKE KKGKGDIE IEIEITDIG IETIEIETE
701 TETIEITGE KEKEKEKE KEKEKEKG KEIEIGREG KKGKEKEIE
751 TEMEIGKIE AEIEEKEK KKKKKKKK KKKKKKKK I
11AA SEQUENCE 1.0
ID O8I247 PRELIMINARY; PRT; 206 AA.
AC O8I247;
DT 01-MAR-2003 (Tremblrel. 23, Created)
DT 01-MAR-2003 (Tremblrel. 23, Last sequence update)
DT 01-MAR-2003 (Tremblrel. 23, Last annotation update)
DE Hypothetical protein.
OS Plasmodium falciparum (isolate 3D7).
OC Eukaryota; Alveolata; Apicomplexa; Haemosporida; Plasmodium.
OX NCBI_TaxID=36329;
RN
RP SEQUENCE FROM N.A.
RC MEDLINE=2255708; PubMed=1236867;
RA Hall N., Pain A., Barriman M., Churcher C., Harris B., Harris D.,
RA Mungall K., Bowman S., Atkin R., Baker S., Barron A., Brooks K.,
RA Buckee C.O., Burrows C., Cherevach I., Chillingworth C.,
RA Chillingworth T., Christodoulou Z., Clark L., Clark C., Corton C.,
RA Cronin A., Davies R., Davis P., Dear P., Darden F., Doggett J.,
RA Fretwell T., Goble A., Goodhead I., Gwilliam R., Hamlin N., Hance Z.,
RA Harper D., Hanger H., Hornsby T., Holroyd S., Horrocks P.,
RA Humphray S., Jagels K., James K.D., Johnson D., Kerhrou A.,
RA Knights A., Konfortov B., Kyes S., Larke N., Lawson D., Leonard N.,
RA Line A., Maddison M., McLean J., Mooney P., Moule S., Murphy L.,
RA Oliver K., Ormond D., Price C., Quail M.A., Rabbittowtsch E.,
RA Rajandream M.A., Rutter S., Rutherford K.M., Sanders M., Simmonds M.,
RA Seeger K., Sharp S., Smith R., Squares R., Squares S., Stevens K.,
RA Taylor K., Tivey A., Unwin L., Whitehead S., Woodward J.,
RA Sulston J.E., Craig A., Newbold C., Barrell B.G.;
RT "Sequence of Plasmodium falciparum chromosomes 1, 3-9 and 13.";
RL Nature 419:527-531 (2002).
DR EMBL; AL031745; CAD49055.1; -.

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KW Hypothetical protein.
SQ SEQUENCE 206 AA; 25047 MW; 1192E49A3DC4523F CRC64;
Q81247 Length: 206 January 30, 2004 08:18 Type: P Check: 5951 ..

1 MEGQHEKNT KIKSKKPLV VSNRKPFPNV EKSKAKPLV RDPSPDSG
51 SNNAPFNA YKPLYDSREQ EKKIEKKLK SKNITQEKD ELKKYNDYK
101 STDILKKKE EERLKAEVL KQEKONILTK NKKPYYSOR KIKKIQEKL
151 SSYSLKAVI KKEKTIQKE RKNIKPTKK KIFLKKKKK KKKKKKKKK
201 KKKKK

11AA SEQUENCE 1.0
ID Q95LV6 PRELIMINARY; PRT; 531 AA.
AC Q95LV6;
DT 01-DEC-2001 (Tremblrel. 19, Created)
DT 01-DEC-2001 (Tremblrel. 19, Last sequence update)
DT 01-OCT-2002 (Tremblrel. 22, Last annotation update)
DE Hypothetical 61.4 kDa protein (fragment). (Cynomolgus monkey).
OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopitheciidae;
OC Cercopitheciinae; Macaca.
OC NCBI_Taxid=9541;
RN [1]
RP SEQUENCE FROM N.A.
RC Tissue=Testis;
RA Hashimoto K., Osada N., Hida M., Kusuda J., Tanuma R., Hirai M.,
RA Terao K., Sugano S.;
RT "Isolation of novel full-length cDNA clones from macaque testis cDNA
RT libraries."
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AB071085; BAB64479.1; -.
KW Hypothetical protein.
FT NON TER 531 531
SQ SEQUENCE 531 AA; 61389 MW; B5596B4F5CDD60C CRC64;
Q95LV6 Length: 531 January 30, 2004 08:18 Type: P Check: 7515 ..

1 MESESSNAN MNVQHEREDK NIOQLPEV PCYSQHLSPS TYOMKDPDC
51 KSRSEPKSE GRSSWNLSTI VQKTEQTHF RESVLEPIG YMKKSPHMQ
101 EGICVGVGK TSFPTGKSE IGSMPHDPW DENPRKWD SISEKTAMP
151 KNLQTVLKL DSSLSMSEY ESRSYTLEFI GKKSITSPKH VTLKTKQLPI
201 SOLPPIRCS TENHRKKQH CFKYMKGRO WYTSIGEAR SATVAKSP
251 SKSMIDKLF NTAAGTILN RTHQNVYGH TTEEKEVGE NVAASLQPL
301 DFFMVLSDS KNQNTTIRLS ERKTIILPKC LTMKEKSP I SQIRKINHF
351 TTKHKKKLES NLKTKKAMW QGENVDTTP NITSFPDPS DIKQSRQOT
401 BIDMRISGLS HTQPTQIESL AEGIARCS DK RSTNLVKT KLHDSSEGEK
451 KOEHLTGMP FYAENFMTNT HLRKDPHLGK SEDVLLGEPF ISKQFYKGN
501 SKKKKKKKK KKKKKKKKK KKKKKKKKK K

11AA SEQUENCE 1.0
ID Q9LXR2 PRELIMINARY; PRT; 517 AA.
AC Q9LXR2;
DT 01-OCT-2000 (Tremblrel. 15, Created)
DT 01-OCT-2000 (Tremblrel. 15, Last sequence update)
DT 01-MAR-2003 (Tremblrel. 23, Last annotation update)
DE Hypothetical 59.7 kDa protein.
GN T20N10.250.
KW Arabidopsis thaliana (Mouse-ear cress).

OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsids.
OC NCBI_Taxid=3702;
RN [1]
RP SEQUENCE FROM N.A.
RA D'Angelo M., Vezzi A., Modesto D., Pigazzi M., Valle G., Mewes H.W.,
RA Rugg S., Lemcke K., Mayer K.F.X., Queller F., Salanoubat M.;
RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RA EU Arabidopsis sequencing project;
RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
DR EMBL; AL353032; CAB88307.1; -.
DR InterPro; IPR001810; F-box.
DR InterPro; IPR005566; FBD.
DR Pfam; PF00646; F-box; 1.
DR SMART; SM00579; FBD; 1.
DR SMART; SM00256; FBOX; 1.
DR PROSITE; PS50181; FBOX; 1.
KW Hypothetical protein.
SQ SEQUENCE 517 AA; 59689 MW; EC6D957D01F86E70 CRC64;
Q9LXR2 Length: 517 January 30, 2004 08:18 Type: P Check: 4143 ..

1 MDLPSLENE LKHILSFLS TKEALTSVL SKRWNLPAF VPYLEPDSV
51 FLHPEERKRE KEGILQSFMD FVDRLVDLHG DSLITFSLK CKTVSDBDHV
101 DRWICNVLAR GVSDDLDFID FDLVSLPHE VGSRTLVVL RVGSESDLYW
151 WQFLCLPML KTLVLDSCWL CIGQFILL ACPALDELDM TNRWKDSNV
201 TVSSSILKEL TIDLHGCSV VNLKSLSPDA PSLYFYICD SLADYQVN
251 LKNLVEAQIN LLLTQAIQEQ VBALNNMLV ADVVPEGLGN AKMLITGLRN
301 VQVLSPDT LEVLSRCCEG MPVFNNLVYL SIWSDMNGW QAMPVLLRNC
351 PHLETILIEG LHAHTDKG DVCDCISRDY KDHSLTSPV KKLQIYFERG
401 TIRELEMIKH FLKIFPLKE MDIYAHNSH TLFKQPTFE RVGKKKKKK
451 KKKKKKKKK KKKKKKIRLN FKPVNKTQEF LKRLADKLCF IPQCLEFLDV
501 DSSLGELLAL AMSDRPS

11AA SEQUENCE 1.0
ID Q8S7D3 PRELIMINARY; PRT; 80 AA.
AC Q8S7D3;
DT 01-JUN-2002 (Tremblrel. 21, Created)
DT 01-JUN-2002 (Tremblrel. 21, Last sequence update)
DT 01-OCT-2002 (Tremblrel. 22, Last annotation update)
DE Hypothetical 9.4 kDa protein.
GN OSJNBA0057121.23.
OS Oryza sativa (Rice).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzaceae; Oryza.
OC NCBI_Taxid=4530;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Nipponbare;
RA Buehl C.R., Yuan Q., Ouyang S., Liu J., Mofiat K.S., Hill J.N.,
RA Gansberger K., Brenner M., Burgess S., Hance M., Shvartsbeyn M.,
RA Tsilirtin T., Riggs F., Hsiao J., Zismann V., Blunt S., Pai G.,
RA Vanabek S.E., Utterback T.R., Feldblum T.V., Kalb E., Quackenbush J.,
RA Salzberg S.L., White O., Fraser C.M.;
RT "Oryza sativa chromosome 10 BAC OSJNBA0057121 genomic sequence.";
RL Submitted (FEB-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC087599; AAL79706.1; -.
DR Gramene; Q8S7D3; -.
KW Hypothetical protein.

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50 SEQUENCE 80 AA; 9362 MW; 0177C863133B21D8 CRC64;

0857D3 Length: 80 January 30, 2004 08:18 Type: P Check: 6479 ..

1 MAVTKEQPIVG RRMSEAAGGA TPGSRMGHW WPATAITRVF LSVIHLDDKK

51 KKKKKKKKK KKKKKKKKK KKKKKKKLLGE

11AA_SEQUENCE 1.0 PRELIMINARY; PRT; 113 AA.

AC Q9L0P6; PRELIMINARY; PRT; 113 AA.

DT 01-OCT-2002 (TREMBLrel. 22, Created)

DT 01-OCT-2002 (TREMBLrel. 22, Last sequence update)

DE 01-MAR-2003 (TREMBLrel. 23, Last annotation update)

GN OJ117_G01.13

OS Oryza sativa (japonica cultivar-group).

OC Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;

OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;

OC Eriarctoidae; Oryzaceae; Oryza.

OX NCBI_TaxID=39947;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=cv. Nipponbare;

RA Sasaki T., Matsumoto T., Yamamoto K.;

RT "Oryza sativa nipponbare(GA3) genomic DNA, chromosome 1, BAC

clone:OJ117_G01.13";

RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.

DR EMBL; AP003374; BAB9330.1; ..

DR Gramene; O8L0P6; ..

DR SEQUENCE 113 AA; 13660 MW; 597DB0EDB2AA3EF CRC64;

Q8L0P6 Length: 113 January 30, 2004 08:18 Type: P Check: 3239 ..

1 MATSIEHMK KKKKKKKKK KKKKKKKRRE REEDDEEEBE

51 EEEELKLNKY IWDIYEAKG INEKLMPGIY VYLAHDECV RLRTGVVVK

101 HEAMEICLFP VQV

11AA_SEQUENCE 1.0 PRELIMINARY; PRT; 260 AA.

AC Q9L6Z9; PRELIMINARY; PRT; 260 AA.

DT 01-OCT-2000 (TREMBLrel. 15, Created)

DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)

DE 01-OCT-2002 (TREMBLrel. 22, Last annotation update)

OS Arabidopsis thaliana (Mouse-ear cress).

OC Arabidops; Viridiplantae; Streptophyta; Tracheophyta;

OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae;

OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.

OX NCBI_TaxID=3702;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=Columbia;

RA Nakamura Y.;

RT "Structural Analysis of Arabidopsis thaliana Chromosome 3. III."

DR EMBL; AP004460; BAA97098.1; ..

DR InterPro; IPR005819; Histone_H5.

DR PRINTS; PR00624; HISTONEH5.

DR SEQUENCE 260 AA; 33307 MW; 43E2394CB8131143 CRC64;

Q9L6Z9 Length: 260 January 30, 2004 08:18 Type: P Check: 6094 ..

1 MDRCIIRKKK KKKKKKKKK KKKKKKKKK KKKKKKKKK

51 KKKKKKKKK KKKKKKKKK KKKKKKKKK KKKKKKKKK

101 KKKKKKKKK KKKKKKKKK KKKKKKKKK KKKKKKKKK

151 KKKKKKKKK KKKKKKKKK KKKKKKKKK KKKKKKKKK

201 KKKKKKKKK KKKKKKKKK KKKKKKKKK KKKKKKKKK KKKKKKKKK

251 INWMGFIYF

11AA_SEQUENCE 1.0 PRELIMINARY; PRT; 215 AA.

AC Q64075; PRELIMINARY; PRT; 215 AA.

DT 01-NOV-1996 (TREMBLrel. 01, Created)

DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)

DE 01-OCT-2002 (TREMBLrel. 22, Last annotation update)

DE Nucleoporin p62 homolog protein (fragment).

OS Rattus sp.

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

OX NCBI_TaxID=10118;

RN [1]

RP SEQUENCE FROM N.A.

RC MEDLINE=95151924; PubMed=7849178;

RA Wang Z.Q., Akmal K.M., Kim K.H.;

RT "An unusual nucleoporin-related messenger ribonucleic acid is present

in the germ cells of rat testis.";

RL Biol. Reprod. 51:1022-1030(1994).

DR EMBL; S75997; AAB3384.1; ..

DR Porph.

DR NON TER

DR SEQUENCE 215 AA; 24593 MW; 098251C97A8FBD88 CRC64;

Q64075 Length: 215 January 30, 2004 08:18 Type: P Check: 8048 ..

1 SGRATSSGD EDCUSSLPF SLSPYKDC EFLERKKKK KKKKKKKKK

51 KKKKKKTGDN AKSVSRQYSL KYTKLEHAE QAKVELDFIL SQKLEEDLL

101 SPLESYKQ SGTIYLQHAD EEREKTYKLA ENIDQOLKRM AQLKDIIIEH

151 INMAGPADT SDPLQIQICKI LNAHMSLOW VDQSSALLQR RVEASRVCE

201 SRKQERSL RIAPD

11AA_SEQUENCE 1.0 PRELIMINARY; PRT; 169 AA.

AC Q9D5G1; PRELIMINARY; PRT; 169 AA.

DT 01-JUN-2001 (TREMBLrel. 17, Created)

DT 01-MAR-2003 (TREMBLrel. 23, Last sequence update)

DE 01-MAR-2003 (TREMBLrel. 23, Last annotation update)

DE Adult male testis cDNA, RIKEN full-length enriched library,

clone:493044P10 product:hypothetical Arginine-rich region containing

protein, full insert sequence (Fragment).

OS Mus musculus (Mouse).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

OX NCBI_TaxID=10090;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=C57BL/6J; TISSUE=Testis;

RA Adachi J., Aizawa K., Akahira S., Akimura T., Arai A., Bono H.,

Arakawa T., Bono H., Carninci P., Fukuda S., Fukunishi Y., Furuno M.,

Hamagaki T., Hara A., Hayatsu N., Hiramoto K., Hiraoka T., Hori F.,

Imotani K., Ishii Y., Itoh M., Izawa M., Kasukawa T., Kato H.,

Kawai J., Kojima Y., Kono H., Kouda M., Koya S., Kurihara C.,

Matsuyama T., Miyazaki A., Nishi K., Nomura K., Numazaki R., Ono M.,

Okazaki Y., Okido T., Owa C., Saito H., Saito R., Sakai C., Sakai K.,

Sano H., Sasaki D., Shibata K., Shibata Y., Shinagawa A., Shiraki T.,

Sogabe Y., Suzuki D., Tagami M., Tagawa A., Takahashi F., Tanaka T.,

Tejima Y., Toya T., Yamamura T., Yasunishi A., Yoshida K., Yoshino M.,

Muramatsu M., Hayashizaki Y.;

RL Submitted (JUN-2000) to the EMBL/GenBank/DBJ databases.

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=C57BL/6J; TISSUE=Testis;

RE MEDLINE=22354683; PubMed=12466851;

RA The FANTOM Consortium,

the RIKEN Genome Exploration Research Group Phase I & II Team;

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RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs."
RL Nature 420:563-573(2002).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Testis;
RX MEDLINE=21085660; PubMed=11217851;
RA RIKEN FANTOM Consortium;
RT "Functional annotation of a full-length mouse cDNA collection."
RL Nature 403:685-690(2001).
RN [4]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Testis;
RX MEDLINE=99279253; PubMed=10349636;
RA Carninci P., Hayashizaki Y.;
RT "High-efficiency full-length cDNA cloning."
RL Meth. Enzymol. 303:19-44(1999).
RN [5]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Testis;
RX MEDLINE=20499374; PubMed=11042159;
RA Carninci P., Shibata Y., Hayatsu N., Sugahara Y., Shibata K., Itoh M.,
RA Komoto H., Okazaki Y., Muramatsu M., Hayashizaki Y.;
RT "Normalization and subtraction of cap-trapper-selected cDNAs to
RT prepare full-length cDNA libraries for rapid discovery of new genes."
RL Genome Res. 10:1617-1630(2000).
RN [6]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Testis;
RX MEDLINE=20530913; PubMed=11076861;
RA Shibata K., Itoh M., Aizawa K., Nagaoka S., Sasaki N., Carninci P.,
RA Komoto H., Akiyama J., Nishi K., Katsunai T., Tashiro H., Itoh M.,
RA Suni N., Ishi Y., Nakamura S., Hazama M., Nishine T., Harada A.,
RA Yamamoto R., Matsumoto H., Sakaguchi S., Ikegami T., Kashiwagi K.,
RA Fujiwara S., Inoue K., Togawa Y., Izawa M., Ohara E., Watanuki M.,
RA Yoneda Y., Ishikawa T., Ozawa K., Tanaka T., Matsura S., Kawai J.,
RA Okazaki Y., Muramatsu M., Inoue Y., Kira A., Hayashizaki Y.;
RT "RIKEN integrated sequence analysis (RISA) system-384-format
RT sequencing pipeline with 384 multicapillary sequencer."
RL Genome Res. 10:1757-1771(2000).
DR EMBL; AK015384; BAB29822.2; -.
KW Hypothetical protein.
FT NON TER
SQ SEQUENCE 169 AA; 19305 MW; 91B9959380A694CC CRC64;

Q9D5G1 Length: 169 January 30, 2004 08:18 Type: P Check: 9388 ..

1 RCTGQAGPOL RALAGPWPR LAPALLSGR ARNIAGLPAA KHAPDSGAS
51 AARLPAPAPH RGGQPGDAAS LSRELASTHG RRLPHCPPL PLAQTSLVP
101 WYRLKKPIS SVQLGRRRR RRRRRRRRR RRRRRRRRK KEEBEEBVA
151 FLDSLIEPRL TSNLHRRH

!!AA_SEQUENCE 1.0
ID Q9D5G1 PRELIMINARY; PRT; 129 AA.
AC Q35807;
DT 01-JAN-1998 (TREMBLrel. 05, Created)
DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE MICOVASCULAR endothelial differentiation protein 2.
GN MDGZ.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Testis;
RX MEDLINE=9817270E; PubMed=9511718;
RA Proels F., Loser B., Marx M.;
RT "Differential expression of osteopontin, PC4, and CEC5, a novel mRNA

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RT species, during in vitro angiogenesis."
RL Exp. Cell Res. 239:1-10(1998).
DR EMBL; Y08769; CAAT0022.1; -.
DR InterPro; IPR000719; Prot_kinase.
DR Pfam; PF00069; Kinase; 1.
DR ProDom; PD000001; Prot_kinase; 1.
DR PROSITE; PS50011; PROTEIN_KINASE_DOM, 1.
KW ATP-binding; Transferase.
SQ SEQUENCE 129 AA; 15080 MW; 38102272BBE2EDB4 CRC64;

Q35807 Length: 129 January 30, 2004 08:18 Type: P Check: 7510 ..

1 MLKPHIVEL LETYSSDGL YWFFPMGDA DLCEIVKRA DAGFVSEAV
51 ASHWYRQILE ALRYCHDNNI HRDVKPCV LLASKKKKK KKKKKKKKK
101 KKKKKIKWEG RDAFWAIPV KSGGVITQ

!!AA_SEQUENCE 1.0
ID Q35807 PRELIMINARY; PRT; 115 AA.
AC Q8BXG9;
DT 01-MAR-2003 (TREMBLrel. 23, Created)
DT 01-MAR-2003 (TREMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Hypothetical arginine-rich region containing protein (Fragment).
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
RX MEDLINE=22354683; PubMed=12466851;
RA The RIKEN Genome Consortium;
RA the RIKEN Genome Exploration Research Group Phase I & II Team;
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs."
RL Nature 420:563-573(2002).
DR EMBL; AK047167; BAC32979.1; -.
KW Hypothetical protein.
FT NON TER
SQ SEQUENCE 115 AA; 13938 MW; 6C0F7EBD8669CF65 CRC64;

Q8BXG9 Length: 115 January 30, 2004 08:18 Type: P Check: 5434 ..

1 GGRGISRFE VSQYTEKPC LKPKKKKKK IIIIIIIIR RRRRRRRRR
51 RRRRRRRRK ETGSHFVALA SLEHLPPC WNLRYRLPH TQIGFNAIKI
101 SIKDTSLSL AFVK

!!AA_SEQUENCE 1.0
ID Q8BXV2 PRELIMINARY; PRT; 154 AA.
AC Q8BXV2;
DT 01-MAR-2003 (TREMBLrel. 23, Created)
DT 01-MAR-2003 (TREMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE Weakly similar to hypothetical 10.3 kDa protein.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J; TISSUE=Cerebellum;
RX MEDLINE=22354683; PubMed=12466851;
RA The RIKEN Genome Consortium;
RA the RIKEN Genome Exploration Research Group Phase I & II Team;
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs."
RL Nature 420:563-573(2002).
DR EMBL; AK082253; BAC38447.1; -.
KW Hypothetical protein.

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SO SEQUENCE 154 AA; 19596 MW; ABE8158A4839A216 CRC64;
QBHV2 Length: 154 January 30, 2004 08:18 Type: P Check: 8958 ..
1 MVOESRKQSN VGQKTSRRRN VFKEVRFSSR VPRDRERGG ERRRRRRRR
51 RRRRRRRRR RRRRRRRRR KRERERESEH EVSNANKDFG LLMHFCAC
101 DEMGRIFISH QKRNMAFLP SGDIIDRYLS YNWVMSLPS ILYYMLKHC
151 GGCT

! FINDPATTERNS on geneseq: * allowing 0 mismatches
! 1 (R,K){20,20} January 30, 2004 07:18 ..

1 AAP20159 ck: 5750 len: 20 ! Aap20159 Sequence of lysine polymer. 8/2002
(R,K){20,20}
(K){20}
1: KKKKKKKKKKKKKKKKKKK

1 AAP61030 ck: 9157 len: 898 ! Aap61030 Entire coded sequence from plasmid
(R,K){20,20}
(K){20}

873: KNITW KKKKKKKKKKKKKKKKKKK KKKKK

874: NITWK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

875: ITWKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}

876: TWKKK KKKKKKKKKKKKKKKKKKK KKK
(K){20}

877: WKKKK KKKKKKKKKKKKKKKKKKK KK
(K){20}

878: KKKKK KKKKKKKKKKKKKKKKKKK K
(K){20}

879: KKKKK KKKKKKKKKKKKKKKKKKK
(K){20}

1 AAP61056 ck: 2017 len: 899 ! Aap61056 Translation of plasmid PAU157 enc
(R,K){20,20}
(K){20}

873: KNITW KKKKKKKKKKKKKKKKKKK KKKKK

874: NITWK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

875: ITWKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

876: TWKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}

877: WKKKK KKKKKKKKKKKKKKKKKKK KKK
(K){20}

878: KKKKK KKKKKKKKKKKKKKKKKKK KK
(K){20}

879: KKKKK KKKKKKKKKKKKKKKKKKK K
(K){20}

880: KKKKK KKKKKKKKKKKKKKKKKKK
(K){20}

1 AAP61082 ck: 7915 len: 898 ! Aap61082 Complete translation of plasmid pA
(R,K){20,20}
(K){20}

873: KNITW KKKKKKKKKKKKKKKKKKK KKKKK

874: NITWK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

875: ITWKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}

876: TWKKK KKKKKKKKKKKKKKKKKKK KKK
(K){20}

877: WKKKK KKKKKKKKKKKKKKKKKKK KK
(K){20}

878: KKKKK KKKKKKKKKKKKKKKKKKK K
(K){20}

879: KKKKK KKKKKKKKKKKKKKKKKKK
(K){20}

1 AAR29580 ck: 4341 len: 657 ! Aar29580 FMR-1 gene product. 3/2003
(R,K){20,20}
(R){20}

19: RRRRP RRRRRRRRRRRRRRRRRR RRRRL

20: RRRPR RRRRRRRRRRRRRRRRRR RRRLG
(R){20}

21: RRRPR RRRRRRRRRRRRRRRRRR RRLGL
(R){20}

22: RPRRR RRRRRRRRRRRRRRRRRR RLGLR
(R){20}

23: PRRRR RRRRRRRRRRRRRRRRRR LGLER
(R){20}

1 AAW03642 ck: 9623 len: 116 ! Aaw03642 Human cannabinoid GPR N-terminal s
(R,K){20,20}
(K){20}

34: QYEDI KKKKKKKKKKKKKKKKKKK KSPFQ
(K){20}

35: YEDIK KKKKKKKKKKKKKKKKKKK SPFOE
(K){20}

1 AAW38839 ck: 801 len: 28 ! Aaw38839 Delivery peptide used in peptide m
(R,K){20,20}
(K){20}

1: KKKKKKKKKKKKKKKKKKK KKKKK

2: K KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

3: KK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

4: KKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

5: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}

6: KKKKK KKKKKKKKKKKKKKKKKKK KKK
(K){20}

7: KKKKK KKKKKKKKKKKKKKKKKKK XK
(K){20}

AAW38840 ck: 2989 len: 29 ! Aaw38840 Delivery peptide used in peptide m

1

(R,K){20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
7: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
8: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}

AAW38841 ck: 5252 len: 30 ! Aaw38841 Delivery peptide used in peptide m

1

(R,K){20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
7: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
8: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
9: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}

AAW38842 ck: 7590 len: 31 ! Aaw38842 Delivery peptide used in peptide m

1

(R,K){20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}

5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKK

6: KKKKK (K){20} KKKKKKKKKKKKKKKKKKKKKKKKK

7: KKKKK (K){20} KKKKKKKKKKKKKKKKKKKKKKKKK

8: KKKKK (K){20} KKKKKKKKKKKKKKKKKKKKKKKKK

9: KKKKK (K){20} KKKKKKKKKKKKKKKKKKKKKKKKK

10: KKKKK (K){20} KKKKKKKKKKKKKKKKKKKKKKKKK

AAW38877 ck: 1129 len: 23 ! Aaw38877 Delivery peptide used in peptide m

1

(R,K){20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}

AAW38843 ck: 3 len: 32 ! Aaw38843 Delivery peptide used in peptide m

1

(R,K){20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
7: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
8: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
9: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
10: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
11: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}

AAW38878 ck: 2949 len: 24 ! Aaw38878 Delivery peptide used in peptide m

1

(R,K){20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20} KVTK
2: K KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20} VTK

AAW38844 ck: 2491 len: 33 ! Aaw38844 Delivery peptide used in peptide m


```

1      (R,K){20,20}
1:      KKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
2:      K KKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
3:      KK KKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
4:      KKK KKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
5:      KKKK KKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
6:      KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
7:      KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
8:      KKKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
9:      KKKKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
10:     KKKKKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
11:     KKKKKKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
12:     KKKKKKKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}

AAW38679 ck: 4644 len: 25 ! Aaw38679 Delivery peptide used in peptide m
1      (R,K){20,20}
1:      KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
2:      K KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
3:      KK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
4:      KKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
5:      KKKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
6:      KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}

AAW38645 ck: 5054 len: 34 ! Aaw38645 Delivery peptide used in peptide m
1      (R,K){20,20}
1:      KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
2:      K KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
3:      KK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
4:      KKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
5:      KKKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}
6:      KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK
      (K){20}

```

```

7: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}

AAW38880 ck: 6814 len: 26 ! Aaw38880 Delivery peptide used in peptide m-
1: KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (R,K) {20,20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}

AAW38846 ck: 7692 len: 35 ! Aaw38846 Delivery peptide used in peptide m-
1: KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (R,K) {20,20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}

```

```
12: KKKK (K){20} KKK
13: KKKK (K){20} KKK
14: KKKK (K){20} KKK

AAW38833 ck: 9248 len: 22 ! Aaw38833 Delivery peptide used in peptide n
(R,K){20,20}
(K){20}
KKKKKKKKKKKKKKKKKKKK KX

1
AAW38834 ck: 986 len: 23 ! Aaw38834 Delivery peptide used in peptide n
(R,K){20,20}
(K){20}
KKKKKKKKKKKKKKKKKKKK KX

1
AAW38835 ck: 2799 len: 24 ! Aaw38835 Delivery peptide used in peptide n
(R,K){20,20}
(K){20}
KKKKKKKKKKKKKKKKKKKK KX

1
AAW38836 ck: 4687 len: 25 ! Aaw38836 Delivery peptide used in peptide n
(R,K){20,20}
(K){20}
KKKKKKKKKKKKKKKKKKKK KX

1
AAW38837 ck: 6650 len: 26 ! Aaw38837 Delivery peptide used in peptide n
(R,K){20,20}
(K){20}
KKKKKKKKKKKKKKKKKKKK KX
1: KKKKKKKKKKKKKKKKKKKKK KX
2: KKKKKKKKKKKKKKKKKKKKK KX
3: KKKKKKKKKKKKKKKKKKKKK KX
4: KKKKKKKKKKKKKKKKKKKKK KX
(K){20}

1
AAW38838 ck: 8688 len: 27 ! Aaw38838 Delivery peptide used in peptide n
(R,K){20,20}
(K){20}
KKKKKKKKKKKKKKKKKKKK KX
1: KKKKKKKKKKKKKKKKKKKKK KX
2: KKKKKKKKKKKKKKKKKKKKK KX
3: KKKKKKKKKKKKKKKKKKKKK KX
4: KKKKKKKKKKKKKKKKKKKKK KX
5: KKKKKKKKKKKKKKKKKKKKK KX
6: KKKKKKKKKKKKKKKKKKKKK KX
(K){20}

1
AAW38796 ck: 9227 len: 22 ! Aaw38796 Delivery peptide used in peptide n
(R,K){20,20}
(K){20}
KKKKKKKKKKKKKKKKKKKK WK

1
AAW38797 ck: 964 len: 23 ! Aaw38797 Delivery peptide used in peptide n
(R,K){20,20}
(K){20}
KKKKKKKKKKKKKKKKKKKK WK
1: KKKKKKKKKKKKKKKKKKKKK WK
2: KKKKKKKKKKKKKKKKKKKKK WK
(K){20}

1
AAW38798 ck: 2776 len: 24 ! Aaw38798 Delivery peptide used in peptide n
(R,K){20,20}
(K){20}
KKKKKKKKKKKKKKKKKKKK KX
1: KKKKKKKKKKKKKKKKKKKKK KX
2: KKKKKKKKKKKKKKKKKKKKK KX
3: KKKKKKKKKKKKKKKKKKKKK KX
(K){20}

1
AAW38799 ck: 4663 len: 25 ! Aaw38799 Delivery peptide used in peptide n
(R,K){20,20}
(K){20}
KKKKKKKKKKKKKKKKKKKK KX
1: KKKKKKKKKKKKKKKKKKKKK KX
2: KKKKKKKKKKKKKKKKKKKKK KX
3: KKKKKKKKKKKKKKKKKKKKK KX
4: KKKKKKKKKKKKKKKKKKKKK KX
(K){20}

1
AAW38800 ck: 6625 len: 26 ! Aaw38800 Delivery peptide used in peptide n
(R,K){20,20}
(K){20}
KKKKKKKKKKKKKKKKKKKK KX
1: KKKKKKKKKKKKKKKKKKKKK KX
2: KKKKKKKKKKKKKKKKKKKKK KX
3: KKKKKKKKKKKKKKKKKKKKK KX
4: KKKKKKKKKKKKKKKKKKKKK KX
(K){20}
```

```
1
(R,K){20,20}
(K){20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
2: K KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKK WK

AAW38801 ck: 8662 len: 27 ! Aaw38801 Delivery peptide used in peptide m
(R,K){20,20}
(K){20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
6: KKKKK KKKKKKKKKKKKKKKKKKK WK

AAW38802 ck: 774 len: 28 ! Aaw38802 Delivery peptide used in peptide m
(R,K){20,20}
(K){20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
6: KKKKK KKKKKKKKKKKKKKKKKKK WK

AAW38803 ck: 2961 len: 29 ! Aaw38803 Delivery peptide used in peptide m
(R,K){20,20}
(K){20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKK KKKKK
```

```
1
(K){20}
3: KK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
6: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
7: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
8: KKKKK KKKKKKKKKKKKKKKKKKK WK

AAW38804 ck: 5223 len: 30 ! Aaw38804 Delivery peptide used in peptide m
(R,K){20,20}
(K){20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
6: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
7: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
8: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
9: KKKKK KKKKKKKKKKKKKKKKKKK WK

AAW38805 ck: 7560 len: 31 ! Aaw38805 Delivery peptide used in peptide m
(R,K){20,20}
(K){20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
6: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
7: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
```

8: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKK
(K) { 20}

9: KKKKK KKKKKKKKKKKKKKKKKKKKK KKK
(K) { 20}

10: KKKKK KKKKKKKKKKKKKKKKKKKKK KKK
(K) { 20}

```
AAW38806 ck: 9972 len: 32 | Aaw38806 Delivery peptide used in peptide m
```

[illegible]

AAW38807 ck: 2459 len: 33 ! Aaw38807 Delivery peptide used in peptide m

[illegible]

```

8: KKKK KKKKKKKKKKKKKKKKKKK KKKK
9: KKKK KKKKKKKKKKKKKKKKKKK KKKK
   (K) {20}
10: KKKK KKKKKKKKKKKKKKKKKKK KKMK
    (K) {20}
11: KKKK KKKKKKKKKKKKKKKKKKK KMK
    (K) {20}
12: KKKK KKKKKKKKKKKKKKKKKKK WK
    (K) {20}

```

AAW38808 ck: 5021 len: 34 | Aaw38808 Delivery peptide used in peptide m

[illegible]

AAW38881	ck: 8859	len: 27	! Aaw38881	Delivery peptide used in peptide m
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[illegible]

5: KKKK (K) {20}
KKKKKKKKKKKKKKKKKKKK VTK

AAW38847 ck: 405 len: 36 ! Aaw38847 Delivery peptide used in peptide m

1

1: (R,K) {20,20}
KKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
2: K KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: KK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKK KKK
(K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKK XK
(K) {20}

AAW38882 ck: 979 len: 28 ! Aaw38882 Delivery peptide used in peptide m

1

(R,K) {20,20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKK KKKVT
(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKK KKVTK
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKK KVTK
(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKK VTK
(K) {20}

1

AAW38848 ck: 3193 len: 37 ! Aaw38848 Delivery peptide used in peptide m

1: (R,K) {20,20}
KKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKK
(K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKK KKK
(K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKKKK XK
(K) {20}

AAW38883 ck: 3174 len: 29 ! Aaw38883 Delivery peptide used in peptide m

1

(R,K) {20,20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKK KKKKV
(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKK KKVVT
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKK KKVTK
(K) {20}

1

6: KKKKK KKKKKKKKKKKKKKKKKKK KVTK
(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKK VTK

AAW38849 ck: 6056 len: 38 | Aaw38849 Delivery peptide used in peptide m

(R,K) {20,20}
(K) {20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
17: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

1

AAW38884 ck: 5444 len: 30 | Aaw38884 Delivery peptide used in peptide m

(R,K) {20,20}
(K) {20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKKKK

1

(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKKV
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKKV

(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKV

(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKV

(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKV

AAW38850 ck: 8994 len: 39 | Aaw38850 Delivery peptide used in peptide m

(R,K) {20,20}
(K) {20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
17: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}
18: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

```
1
AAW3885 ck: 7789 len: 31 ! Aaw3885 Delivery peptide used in peptide m
(R,K){20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
AAW3851 ck: 2007 len: 40 ! Aaw3851 Delivery peptide used in peptide m
(R,K){20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
12: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
```

```
1
14: KKKKK (K){20} KKKKK
(K){20}
15: KKKKK (K){20} KKKKK
(K){20}
16: KKKKK (K){20} KKKKK
(K){20}
17: KKKKK (K){20} KKKKK
(K){20}
18: KKKKK (K){20} KKK
(K){20}
19: KKKKK (K){20} KKK
(K){20}
AAW3886 ck: 209 len: 32 ! Aaw3886 Delivery peptide used in peptide m
(R,K){20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKKKK
(K){20}
AAW3852 ck: 5095 len: 41 ! Aaw3852 Delivery peptide used in peptide m
(R,K){20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
```

6: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
17: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
18: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
19: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
20: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

AAW38887 ck: 2704 len: 33 1 Aaw38887 Delivery peptide used in peptide m

1: (R, K) {20, 20}
(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

9: KKKKK KKKKKKKKKKKKKKKKKKKKK KKVTK
(K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKK KVTKK
(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKK VTK
(K) {20}

AAW38853 ck: 8258 len: 42 1 Aaw38853 Delivery peptide used in peptide m

1: (R, K) {20, 20}
(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
17: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
18: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
19: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
20: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

21: KKKK KKKKKKKKKKKKKKKKKKK KK

AAW3888 ck: 5274 len: 34 ! Aaw3888 Delivery peptide used in peptide m

(R,K) {20,20}

(K) {20}

1: KKKKKKKKKKKKKKKKKKKKKKKKKKK

2: K KKKKKKKKKKKKKKKKKKKKKKKKK

3: KK KKKKKKKKKKKKKKKKKKKKKKKKK

4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK

5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKK

6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

7: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

8: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

9: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

10: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

11: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

12: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

AAW38854 ck: 1496 len: 43 ! Aaw38854 Delivery peptide used in peptide m

(R,K) {20,20}

(K) {20}

1: KKKKKKKKKKKKKKKKKKKKKKKKKKK

2: K KKKKKKKKKKKKKKKKKKKKKKKKKKK

3: KK KKKKKKKKKKKKKKKKKKKKKKKKKKK

4: KKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

7: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

8: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

9: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

10: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

11: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

12: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

13: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

14: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

15: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

16: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

17: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

18: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

19: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

20: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

21: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

22: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKKKKKKKK

AAW38889 ck: 7919 len: 35 ! Aaw38889 Delivery peptide used in peptide m

(R,K) {20,20}

(K) {20}

1: KKKKKKKKKKKKKKKKKKKKKKKKKKK

2: K KKKKKKKKKKKKKKKKKKKKKKKKKKK

3: KK KKKKKKKKKKKKKKKKKKKKKKKKKKK

4: KKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

7: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

8: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

9: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

10: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

11: KKKK (K) {20}
12: KKKK (K) {20}
13: KKKK (K) {20}

AAW38890 ck: 639 len: 36 ! Aaw38890 Delivery peptide used in peptide m

1 (R, K) {20, 20}

1: KKKK (K) {20}
2: K KKKK (K) {20}
3: KK KKKK (K) {20}
4: KK KKKK (K) {20}
5: KKK KKKK (K) {20}
6: KKKK KKKK (K) {20}
7: KKKK KKKK (K) {20}
8: KKKK KKKK (K) {20}
9: KKKK KKKK (K) {20}
10: KKKK KKKK (K) {20}
11: KKKK KKKK (K) {20}
12: KKKK KKKK (K) {20}
13: KKKK KKKK (K) {20}
14: KKKK KKKK (K) {20}

AAW38891 ck: 3434 len: 37 ! Aaw38891 Delivery peptide used in peptide m

1 (R, K) {20, 20}

1: KKKK (K) {20}
2: K KKKK (K) {20}
3: KK KKKK (K) {20}
4: KK KKKK (K) {20}
5: KKK KKKK (K) {20}

6: KKKK (K) {20}
7: KKKK (K) {20}
8: KKKK (K) {20}
9: KKKK (K) {20}
10: KKKK (K) {20}
11: KKKK (K) {20}
12: KKKK (K) {20}
13: KKKK (K) {20}
14: KKKK (K) {20}
15: KKKK (K) {20}

AAW38892 ck: 6304 len: 38 ! Aaw38892 Delivery peptide used in peptide m

1 (R, K) {20, 20}

1: KKKK (K) {20}
2: K KKKK (K) {20}
3: KK KKKK (K) {20}
4: KK KKKK (K) {20}
5: KKK KKKK (K) {20}
6: KKKK KKKK (K) {20}
7: KKKK KKKK (K) {20}
8: KKKK KKKK (K) {20}
9: KKKK KKKK (K) {20}
10: KKKK KKKK (K) {20}
11: KKKK KKKK (K) {20}
12: KKKK KKKK (K) {20}
13: KKKK KKKK (K) {20}

(K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKK KKVTK
(K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKK KVTK
(K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKK VTK

AAW38893 ck: 9249 len: 39 ! Aaw38893 Delivery peptide used in peptide m

1

(R,K) {20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
17: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK VTK

AAW38894 ck: 2269 len: 40 ! Aaw38894 Delivery peptide used in peptide m

1

(R,K) {20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
(K) {20}

2: K KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

3: KK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

7: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

8: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

9: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

10: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

11: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

12: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

13: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

14: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

15: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

16: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

17: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

(K) {20}

18: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK VTK

AAW38895 ck: 5364 len: 41 ! Aaw38895 Delivery peptide used in peptide m

1

(R,K) {20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

1

```
7: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKVT
   (K) {20}
17: KKKKK KKKKKKKKKKKKKKKKKKKKK KKVTK
   (K) {20}
18: KKKKK KKKKKKKKKKKKKKKKKKKKK KVTK
   (K) {20}
19: KKKKK KKKKKKKKKKKKKKKKKKKKK VTK
   (K) {20}

AAW38896 ck: 8534 len: 42 ! Aaw38896 Delivery peptide used in peptide m
(R,K) {20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
```

1

```
11: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKV
   (K) {20}
17: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKVT
   (K) {20}
18: KKKKK KKKKKKKKKKKKKKKKKKKKK KKVTK
   (K) {20}
19: KKKKK KKKKKKKKKKKKKKKKKKKKK KVTK
   (K) {20}
20: KKKKK KKKKKKKKKKKKKKKKKKKKK VTK
   (K) {20}

AAW38897 ck: 1779 len: 43 ! Aaw38897 Delivery peptide used in peptide me
(R,K) {20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
   (K) {20}
```

14: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 15: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 16: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 17: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 18: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 19: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 20: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 21: KKKK KKKKKKKKKKKKKKKKKKK KKKK

AAW38898 ck: 5099 len: 44 ! Aaw38898 Delivery peptide used in peptide m

1

1: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 2: K KKKKKKKKKKKKKKKKKKKKK KKKK
 3: KK KKKKKKKKKKKKKKKKKKKKK KKKK
 4: KK KKKKKKKKKKKKKKKKKKKKK KKKK
 5: KKK KKKKKKKKKKKKKKKKKKKKK KKKK
 6: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 7: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 8: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 9: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 10: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 11: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 12: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 13: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 14: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 15: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK

16: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 17: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 18: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 19: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 20: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 21: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 22: KKKK KKKKKKKKKKKKKKKKKKK KKKK

AAW38809 ck: 7658 len: 35 ! Aaw38809 Delivery peptide used in peptide m

1

1: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 2: K KKKKKKKKKKKKKKKKKKKKK KKKK
 3: KK KKKKKKKKKKKKKKKKKKKKK KKKK
 4: KK KKKKKKKKKKKKKKKKKKKKK KKKK
 5: KKK KKKKKKKKKKKKKKKKKKKKK KKKK
 6: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 7: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 8: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 9: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 10: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 11: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 12: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 13: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK
 14: KKKK KKKKKKKKKKKKKKKKKKKKK KKKK

AAW38810 ck: 370 len: 36 ! Aaw38810 Delivery peptide used in peptide m

1

1: KKKK KKKKKKKKKKKKKKKKKKK KKKK

```
2: K (K) {20}
3: KK (K) {20}
4: KKK (K) {20}
5: KKKK (K) {20}
6: KKKKK (K) {20}
7: KKKKKK (K) {20}
8: KKKKKKK (K) {20}
9: KKKKKKKK (K) {20}
10: KKKKKKKKK (K) {20}
11: KKKKKKKKKK (K) {20}
12: KKKKKKKKKKK (K) {20}
13: KKKKKKKKKKKK (K) {20}
14: KKKKKKKKKKKKK KWK
15: KKKKKKKKKKKKKKKKKKK KWK
```

AAW38811 ck: 3157 len: 37 ! Aaw38811 Delivery peptide used in peptide p

```
1: (R,K) {20,20}
2: K (K) {20}
3: KK (K) {20}
4: KKK (K) {20}
5: KKKK (K) {20}
6: KKKKK (K) {20}
7: KKKKKK (K) {20}
8: KKKKKKK (K) {20}
9: KKKKKKKK (K) {20}
```

```
10: KKKKK (K) {20}
11: KKKKKK (K) {20}
12: KKKKKKK (K) {20}
13: KKKKKKKK (K) {20}
14: KKKKKKKKK (K) {20}
15: KKKKKKKKKK (K) {20}
16: KKKKKKKKKKK KWK
```

AAW38812 ck: 6019 len: 38 ! Aaw38812 Delivery peptide used in peptide m

```
1: (R,K) {20,20}
2: K (K) {20}
3: KK (K) {20}
4: KKK (K) {20}
5: KKKK (K) {20}
6: KKKKK (K) {20}
7: KKKKKK (K) {20}
8: KKKKKKK (K) {20}
9: KKKKKKKK (K) {20}
10: KKKKKKKKK (K) {20}
11: KKKKKKKKKK (K) {20}
12: KKKKKKKKKKK (K) {20}
13: KKKKKKKKKKKK (K) {20}
14: KKKKKKKKKKKKK KWK
15: KKKKKKKKKKKKKKKKKKK KWK
16: KKKKKKKKKKKKKKKKKKKKK KWK
```

17: KKKKK (K) {20}
KKKKKKKKKKKKKKKKKKKK WK

AAW38813 ck: 8956 len: 39 1 Aaw38813 Delivery peptide used in peptide m

1: (R,K) {20,20}
(K) {20}
KKKKKKKKKKKKKKKKKKKK KKKKK

2: K KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

3: KK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

4: KKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

5: KKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

6: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

7: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

8: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

9: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

10: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

11: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

12: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

13: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

14: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

15: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

16: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

17: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

18: KKKKK KKKKKKKKKKKKKKKKKKKKK WK
(K) {20}

AAW38814 ck: 1968 len: 40 1 Aaw38814 Delivery peptide used in peptide m

1: (R,K) {20,20}
(K) {20}
KKKKKKKKKKKKKKKKKKKKKK KKKKK

2: K KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

3: KK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

4: KKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

5: KKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

6: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

7: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

8: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

9: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

10: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

11: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

12: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

13: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

14: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

15: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

16: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

17: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

18: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

19: KKKKK KKKKKKKKKKKKKKKKKKKKK WK
(K) {20}

AAW38815 ck: 5055 len: 41 1 Aaw38815 Delivery peptide used in peptide m

1: (R,K) {20,20}
(K) {20}
KKKKKKKKKKKKKKKKKKKKKK KKKKK

2: K KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

3: KK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

4: KKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

5: KKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

6: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

7: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

[illegible]

```

1
AAW38816  ck: 8217  len: 42  ! Aaw38816  Delivery peptide used in peptide m

1:      (R,K) {20,20}
      (K) {20}
      KKKKKKKKKKKKKKKKKKKKKKKKKKK

2:      (K) {20}
      K KKKKKKKKKKKKKKKKKKKKKKKKKKK

3:      (K) {20}
      KK KKKKKKKKKKKKKKKKKKKKKKKKKKK

4:      (K) {20}
      KKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

5:      (K) {20}
      KKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

6:      (K) {20}
      KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

7:      (K) {20}
      KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

8:      (K) {20}
      KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

9:      (K) {20}
      KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

10:      (K) {20}
      KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

```

11:	KKKKK	KKKKK{20}	KKKKK
12:	KKKKK	(K){20}	KKKKK
13:	KKKKK	(K){20}	KKKKK
14:	KKKKK	(K){20}	KKKKK
15:	KKKKK	(K){20}	KKKKK
16:	KKKKK	(K){20}	KKKKK
17:	KKKKK	(K){20}	KKKKK
18:	KKKKK	(K){20}	KKKKK
19:	KKKKK	(K){20}	KKKKK
20:	KKKKK	(K){20}	KKKKK
21:	KKKKK	(K){20}	KKKKK

```

1                                     AAW38917   ck: 1454   len: 43      | Aaw38917 Delivery peptide used in peptide ma

(R,X){20,20}
1:  KKKKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

2:  K KKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

3:  KK KKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

4:  KKK KKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

5:  KKKK KKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

6:  KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

7:  KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

8:  KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

9:  KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

10: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

11: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

12: KKKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK
    (X){20}

```


13: KKKK (K) {20}
14: KKKK (K) {20}
15: KKKK (K) {20}
16: KKKK (K) {20}
17: KKKK (K) {20}
18: KKKK (K) {20}
19: KKKK (K) {20}
20: KKKK (K) {20}
21: KKKK (K) {20}
22: KKKK (K) {20}

1 AAW24865 ck: 2211 len: 40 ! Aaw24865 Bifunctional peptide I for binding
(R,K) {20,20}
21: YEDES KKKKKKKKKKKKKKKKKKK

1 AAW24450 ck: 8137 len: 45 ! Aaw24450 Nucleic acid (NA) binding peptide
(R,K) {20,20}
4: YKA KKKKKKKKKKKKKKKKKKK
5: YPAK KKKKKKKKKKKKKKKKKKK
6: YPAK KKKKKKKKKKKKKKKKKKK
7: KAKK KKKKKKKKKKKKKKKKKKK
8: AKKK KKKKKKKKKKKKKKKKKKK
9: KKKK KKKKKKKKKKKKKKKKKKK
10: KKKK KKKKKKKKKKKKKKKKKKK
11: KKKK KKKKKKKKKKKKKKKKKKK
12: KKKK KKKKKKKKKKKKKKKKKKK
13: KKKK KKKKKKKKKKKKKKKKKKK
(K) {20}

14: KKKK KKKKKKKKKKKKKKKKKKK
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24: KKKK (K) {20}

1 AAW21590 ck: 4875 len: 30 ! Aaw21590 Antibiotic potentiating peptide #2
(R,K) {20,20}
1: KKKKKKKKKKKKKKKKKKK
2: K KKKKKKKKKKKKKKKKKKK
3: KK KKKKKKKKKKKKKKKKKKK
4: KKK KKKKKKKKKKKKKKKKKKK
5: KKKK KKKKKKKKKKKKKKKKKKK
6: KKKK KKKKKKKKKKKKKKKKKKK
7: KKKK KKKKKKKKKKKKKKKKKKK
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9: KKKK KKKKKKKKKKKKKKKKKKK
10: KKKK KKKKKKKKKKKKKKKKKKK
11: KKKK KKKKKKKKKKKKKKKKKKK
(K) {20}

1 AAW21591 ck: 5075 len: 434 ! Aaw21591 Antibiotic potentiating peptide #3
(R,K) {20,20}

[illegible]

25:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
26:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
27:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
28:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
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30:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
31:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
32:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
33:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
34:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
35:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
36:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
37:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
38:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
39:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
40:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
41:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
42:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
43:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
44:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
45:	XXXXX	XXXXXXXXXXXXXXXXXXXXX	XXXXX
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415: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
AAW65939 ck: 1569 len: 40 ! Aaw65939 Polylysine peptide NBC32. 11/1998
(R,K) {20,20}
2: T KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: TK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
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18: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
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AAW48808 ck: 9658 len: 56 ! Aaw48808 Homo sapiens clone CG109_1 protein
(R,K) {20,20}
34: EEPRE KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
35: EPREK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
36: PREKK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

37: REKK KKKKKKKKKKKKKKKKKKK

1 AAW45801 ck: 9500 len: 39 ! Aaw45801 One chain of a bombesin dimer. 6/11
(R,K){20,20}
(K){20}

2: C KKKKKKKKKKKKKKKKKKK XGGGQ

1 AAW45800 ck: 9056 len: 39 ! Aaw45800 One chain of an alpha-melanocyte s
(R,K){20,20}
(K){20}

18: VGGGX KKKKKKKKKKKKKKKKKKK CX

1 AAW45802 ck: 7664 len: 35 ! Aaw45802 One chain of an alpha-MSH receptor
(R,K){20,20}
(K){20}

14: VGGGX KKKKKKKKKKKKKKKKKKK CX

1 AAY43246 ck: 9752 len: 32 ! Aay43246 Cell-surface molecule binding pept
(R,K){20,20}
(K){20}

12: SGSGS KKKKKKKKKKKKKKKKKKK K

13: GSGSK KKKKKKKKKKKKKKKKKKK

1 AAY07213 ck: 2211 len: 40 ! Aay07213 Peptide transfection vector #1. 7/7
(R,K){20,20}
(K){20}

21: YEDES KKKKKKKKKKKKKKKKKKK

1 AAY12950 ck: 5821 len: 62 ! Aay12950 Amino acid sequence of a human sec
(R,K){20,20}
(K){20}

40: QFQAS KKKKKKKKKKKKKKKKKKK KKK

1 41: FQASK KKKKKKKKKKKKKKKKKKK KK
(K){20}

42: QASKK KKKKKKKKKKKKKKKKKKK K

1 43: ASKKK KKKKKKKKKKKKKKKKKKK
(K){20}

1 AAB59105 ck: 8456 len: 27 ! Aab59105 Breast and ovarian cancer associat
(R,K){20,20}
(K){20}

6: NSAXX KKKKKKKKKKKKKKKKKKK KK

1 7: SAXXK KKKKKKKKKKKKKKKKKKK K
(K){20}

8: AXXKK KKKKKKKKKKKKKKKKKKK

1 AAB53249 ck: 4945 len: 59 ! Aab53249 Human colon cancer antigen protein
(R,K){20,20}

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27: ALISL KKKKKKKKKKKKKKKKKKKKK KKKKK
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28: LLSLK KKKKKKKKKKKKKKKKKKKKK KKKKK
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      (K) {20}
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      (K) {20}

AAB53659 ck: 3850 len: 184 | Aab53659 Human colon cancer antigen protein
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      (K) {20}

AAB53800 ck: 296 len: 69 | Aab53800 Human colon cancer antigen protein
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      (R, K) {20, 20}
      (K) {20}
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      (K) {20}
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      (K) {20}

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 38: KKKKK (K) {20} KKKKG
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AAB53806 ck: 8373 len: 64 i Aab53806 Human colon cancer antigen protein
 (R,K) {20,20}

25: IDCDK KKKKK (K) {20} KKKKK
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 27: CDSKK KKKKK (K) {20} KKKKK
 28: DSKKK KKKKK (K) {20} KKKKK
 29: SKKKK KKKKK (K) {20} KKKKK
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AAB53977 ck: 7756 len: 75 i Aab53977 Human colon cancer antigen protein
 (R,K) {20,20}

24: QFCCL KKKKK (K) {20} KKKKK
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AAB53980 ck: 881 len: 45 i Aab53980 Human colon cancer antigen protein
 (R,K) {20,20}

8: TCSFQ KKKKK (K) {20} KKKKK

9: CSFQK KKKKKKKKKKKKKKKKKKKKK KKKKG
 (K) {20}
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 (K) {20}
 11: FQKKK KKKKKKKKKKKKKKKKKKKKK KGGGR
 (K) {20}
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AAB54314 ck: 6235 len: 55 ! Aab54314 Human pancreatic cancer antigen pr

(R,K) {20,20}

(K) {20}

24: FFXKK KKKKKKKKKKKKKKKKKKKKK KKKKK

25: FXXXX KKKKKKKKKKKKKKKKKKKKK KKKKK

26: XXXKK KKKKKKKKKKKKKKKKKKKKK KKKKK

27: XXXKK KKKKKKKKKKKKKKKKKKKKK KKKKK

28: XKKKK KKKKKKKKKKKKKKKKKKKKK KKKKG

29: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKGG

30: KKKKK KKKKKKKKKKKKKKKKKKKKK KGGGR

31: KKKKK KKKKKKKKKKKKKKKKKKKKK KGGRF

32: KKKKK KKKKKKKKKKKKKKKKKKKKK GGGP

AAB56121 ck: 5941 len: 125 ! Aab56121 Human secreted protein sequence en

(R,K) {20,20}

(K) {20}

87: ILKSQ KKKKKKKKKKKKKKKKKKKKK KKKKK

88: LKSQK KKKKKKKKKKKKKKKKKKKKK KKKKK

89: KSQKK KKKKKKKKKKKKKKKKKKKKK KKKKK

90: SQKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

91: QKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

92: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

93: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

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 (K) {20}

102: KKKKK KKKKKKKKKKKKKKKKKKKKK KGGP
 (K) {20}

103: KKKKK KKKKKKKKKKKKKKKKKKKKK GGP
 (K) {20}

AAB27956 ck: 6732 len: 139 ! Aab27956 Human secreted protein seq ID NO:

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92: FRPGK KKKKKKKKKKKKKKKKKKKKK KKKKK

93: RPKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

94: PKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

95: GKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

96: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

97: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

98: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

99: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

100: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

101: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

102: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

103: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

104: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

105: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

4: /EKK KKKKKKKKKKKKKKKKKKKKK KKKGR
(K) {20}
5: EKKK KKKKKKKKKKKKKKKKKKKKK KKKRP
(K) {20}
6: EKKKK KKKKKKKKKKKKKKKKKKKKK KGRPF
(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKK GRPFK
(K) {20}

AAB44380 ck: 8490 len: 42 ! Aab44380 Human secreted protein encoded by
(R,K) {20,20}
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22: IWKKI KKKKKKKKKKKKKKKKKKKKK K
(K) {20}

23: WKIK KKKKKKKKKKKKKKKKKKKKK
(K) {20}

AAB44331 ck: 8131 len: 66 ! Aab44331 Human secreted protein sequence en
(R,K) {20,20}
(K) {20}

40: IHHPK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

41: HHHPK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

42: HHHPK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

43: KPPKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

44: PKKKK KKKKKKKKKKKKKKKKKKKKK KKK
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45: KKKKK KKKKKKKKKKKKKKKKKKKKK KK
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46: KKKKK KKKKKKKKKKKKKKKKKKKKK K
(K) {20}

47: KKKKK KKKKKKKKKKKKKKKKKKKKK
(K) {20}

AAB23585 ck: 9893 len: 36 ! Aab23585 AbK21 linker peptide. 1/2001
(R,K) {20,20}
(K) {20}

15: PGSGS KKKKKKKKKKKKKKKKKKKKK KG
(K) {20}

16: GSGSK KKKKKKKKKKKKKKKKKKKKK G
(K) {20}

AAB23586 ck: 58 len: 36 ! Aab23586 GeK21 linker peptide. 1/2001
(R,K) {20,20}
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15: SGSGS KKKKKKKKKKKKKKKKKKKKK KG
(K) {20}

16: GSGSK KKKKKKKKKKKKKKKKKKKKK G
(K) {20}

AAB23591 ck: 7684 len: 630 ! Aab23591 Modified fibre protein encoded in
(R,K) {20,20}

596: PGSGS KKKKKKKKKKKKKKKKKKKKK KGSYS
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597: GSGSK KKKKKKKKKKKKKKKKKKKKK GSYSM
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AAB23592 ck: 7647 len: 630 ! Aab23592 Modified fibre protein encoded in
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(K) {20}

596: GSGGS KKKKKKKKKKKKKKKKKKKKK KGSYS
(K) {20}

597: GSGSK KKKKKKKKKKKKKKKKKKKKK GSYSM
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AAB23593 ck: 2249 len: 640 ! Aab23593 Modified fibre protein encoded in
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(K) {20}

596: PGSGS KKKKKKKKKKKKKKKKKKKKK KGSAB
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597: GSGSK KKKKKKKKKKKKKKKKKKKKK GSAEK
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AAB23594 ck: 2612 len: 640 ! Aab23594 Modified fibre protein encoded in
(R,K) {20,20}
(K) {20}

596: SGSGS KKKKKKKKKKKKKKKKKKKKK KGSAB
(K) {20}

597: GSGSK KKKKKKKKKKKKKKKKKKKKK GSAEK
(K) {20}

AAB13780 ck: 7317 len: 21 ! Aab13780 Soluble peptide antigen pK. 11/2001
(R,K) {20,20}
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(K) {20}

AAB13783 ck: 4553 len: 45 ! Aab13783 Soluble tandem pKa/ pK peptide con
(R,K) {20,20}
(K) {20}

26: AAAAA KKKKKKKKKKKKKKKKKKKKK
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AAB13784 ck: 4126 len: 44 ! Aab13784 Soluble tandem HA/ pK peptide conj
(R,K) {20,20}
(K) {20}

25: DGMYG KKKKKKKKKKKKKKKKKKKKK
(K) {20}

AAG00834 ck: 6330 len: 103 ! Aag00834 Human secreted protein, SEQ ID NO:
(R,K) {20,20}
(K) {20}

83: RKRQK KKKKKKKKKKKKKKKKKKKKK K
(K) {20}

84: KRQK KKKKKKKKKKKKKKKKKKKKK
(K) {20}

AAY98493 ck: 8137 len: 45 ! Aay98493 Peptide #5 used in nucleic acid tr
(R,K) {20,20}
(K) {20}

4: YKA KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

[illegible]

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AAV98497 ck: 4925 len: 100 ! Aay98497 Peptide #8 used in nucleic acid tre
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AY81805 ck: 1885 len: 351 1 AY81805 Murine mahogany protein sequence at
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AAV56902 ck: 4875 len: 30 i Aay56902 (Lys)30 protein sequence. 4/2000

1: (R,K) {20,20} KKKK
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AAV56903 ck: 5075 len: 434 i Aay56903 (Lys)434 protein sequence. 4/2000

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AAV86248 ck: 8783 len: 128 1 Aay86248 Human secreted protein HCHPP68,SRK
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AAVS9040 ck: 4361 len: 59 1 AayS9040 Nuclear ligand for transporting nu
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AAVS9042 ck: 4925 len: 100 ! Aavs9042 amino acid polymer seq ID NO: 62 c

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(K) {20}
21: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
22: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
23: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
24: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

ABP66883 ck: 3983 len: 292 ! Abp66883 Human polypeptide SEQ ID NO 604. 1
(R,K) {20,20}
(R,K) {20}
273: QVPAP KKKKKKKKKKKKKKKKKKK

ABP67072 ck: 7611 len: 315 ! Abp67072 Human polypeptide SEQ ID NO 793. 1
(R,K) {20,20}
(R,K) {20}
273: QVPAP KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
274: VFPAP KKKKKKKKKKKKKKKKKKK KKKKK

ABG92583 ck: 7907 len: 39 ! Abg92583 Human DNA-binding protein #9. 11/2
(R,K) {20,20}
(K) {20}
9: YFEDL KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
10: FEDLK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
11: EDLKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
12: DLKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
13: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

ABG92588 ck: 9194 len: 87 ! Abg92588 Human DNA-binding protein #14. 11/
(R,K) {20,20}
(K) {20}
52: KIILL KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
53: IILKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
54: ILKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
55: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

56: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
57: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

ABG92589 ck: 8659 len: 104 ! Abg92589 Human DNA-binding protein #15. 11/
(R,K) {20,20}
(K) {20}
75: PLGGQ KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
76: LGGQK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
77: GGQKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
78: GQKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
79: QKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
80: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

ABG92592 ck: 9398 len: 48 ! Abg92592 Human DNA-binding protein #18. 11/
(R,K) {20,20}
(K) {20}
2: Q KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: QK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
4: QKK KKKKKKKKKKKKKKKKKKK KKKKK

ABG92598 ck: 8278 len: 53 ! Abg92598 Human DNA-binding protein #24. 11/
(R,K) {20,20}
(K) {20}
30: NCGIL KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
31: CGILK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
32: GILKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
33: ILKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
34: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK

ABG92599 ck: 444 len: 66 ! Abg92599 Human DNA-binding protein #25. 11/
(R,K) {20,20}
(K) {20}
35: SWTFS KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
36: MTFSK KKKKKKKKKKKKKKKKKKK KKKKK

ABG92600 ck: 5503 len: 50 ! Abg92600 Human DNA-binding protein #26. 11/
(R,K) {20,20}

30: IICLL (K) {20}
XXXXXXXXXXXXXXXXXXXX K
31: ICLLK (K) {20}
XXXXXXXXXXXXXXXXXXXX

ABG92605 ck: 5691 len: 108 ! Abg92605 Human DNA-binding protein #31. 11/

(R,K) {20,20}
(K) {20}

78: VRPCL XXXXXXXXXXXXXXXXXXXX KKKKX

79: RPCLK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

80: PCLLK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

81: CLKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

82: LKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

83: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

84: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

85: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

86: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKX

87: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KX

88: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX K

89: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX

ABG92613 ck: 6029 len: 63 ! Abg92613 Human DNA-binding protein #39. 11/

(R,K) {20,20}

(K) {20}

40: KLTLL XXXXXXXXXXXXXXXXXXXX ISWG

ABG92621 ck: 7170 len: 63 ! Abg92621 Human DNA-binding protein #47. 11/

(R,K) {20,20}

(K) {20}

37: TPERSA XXXXXXXXXXXXXXXXXXXX KKKKX

38: PSRAK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

39: SRAKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

40: RAKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

41: AKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKX

(K) {20}

42: KKKKK XXXXXXXXXXXXXXXXXXXX KK
(K) {20}
43: KKKKK XXXXXXXXXXXXXXXXXXXX K
44: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX

ABG92625 ck: 6110 len: 61 ! Abg92625 Human DNA-binding protein #51. 11/

(R,K) {20,20}

(K) {20}

28: RPTRP XXXXXXXXXXXXXXXXXXXX KKKKX

29: PTRPK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

30: TRPKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

31: RPKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

32: PKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

33: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKG

34: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKGG

35: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKGG

36: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KGGGG

37: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX GGGGG

ABG92626 ck: 5764 len: 74 ! Abg92626 Human DNA-binding protein #52. 11/

(R,K) {20,20}

(K) {20}

40: EFLSA XXXXXXXXXXXXXXXXXXXX KKKKX

41: FLSSAK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

42: LSAAK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

43: SAKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

44: AKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

45: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

46: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

47: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

48: KKKKK (K) {20}
XXXXXXXXXXXXXXXXXXXX KKKKX

49: KKKKK (K) {20}
50: KKKKK (K) {20}
51: KKKKK (K) {20}
52: KKKKK (K) {20}
53: KKKKK (K) {20}
54: KKKKK (K) {20} X

1
ABG92627 ck: 9217 len: 68 ! Abg92627 Human DNA-binding protein #53. 11/
(R,K) {20,20}
38: FLFPE KKKKKKKKKKKKKKKKKKK (K) {20}
39: LFPEK KKKKKKKKKKKKKKKKKKK (K) {20}
40: FPEKK KKKKKKKKKKKKKKKKKKK (K) {20}
41: PEKKK KKKKKKKKKKKKKKKKKKK (K) {20}
42: EKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
43: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
44: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
45: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} GXXF

1
ABG92629 ck: 8162 len: 79 ! Abg92629 Human DNA-binding protein #55. 11/
(R,K) {20,20}
41: VRPRV RKKKKKKKKKKKKKKKKKK (R,K) {20}
42: RPRVR KKKKKKKKKKKKKKKKKKK (K) {20}
43: PRVRK KKKKKKKKKKKKKKKKKKK (K) {20}
44: RVRKK KKKKKKKKKKKKKKKKKKK (K) {20} KGRF
45: VRKKK KKKKKKKKKKKKKKKKKKK (K) {20} GGRF

1
ABG92658 ck: 285 len: 118 ! Abg92658 Human DNA-binding protein #84. 11/
(R,K) {20,20}
98: EKHKQ KKKKKKKKKKKKKKKKKKK (R,K) {20} G

1
ABG92659 ck: 5509 len: 58 ! Abg92659 Human DNA-binding protein #85. 11/
(R,K) {20,20}
36: FYFVC KKKKKKKKKKKKKKKKKKK (R,K) {20}
37: YFVCK KKKKKKKKKKKKKKKKKKK (K) {20} KK
38: FVCKK KKKKKKKKKKKKKKKKKKK (K) {20} K
39: VCKKK KKKKKKKKKKKKKKKKKKK (K) {20}

1
ABG92660 ck: 9074 len: 66 ! Abg92660 Human DNA-binding protein #86. 11/
(R,K) {20,20}
40: LVQCE KKKKKKKKKKKKKKKKKKK (R,K) {20}
41: VQCEK KKKKKKKKKKKKKKKKKKK (K) {20}
42: QCEKK KKKKKKKKKKKKKKKKKKK (K) {20}
43: CEKKK KKKKKKKKKKKKKKKKKKK (K) {20}
44: EKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKK
45: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KK
46: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} K
47: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}

1
ABG92661 ck: 8528 len: 150 ! Abg92661 Human DNA-binding protein #87. 11/
(R,K) {20,20}
113: SRNTV KKKKKKKKKKKKKKKKKKK (R,K) {20}
114: RNTVK KKKKKKKKKKKKKKKKKKK (K) {20}
115: NTVRK KKKKKKKKKKKKKKKKKKK (K) {20}
116: TVRKK KKKKKKKKKKKKKKKKKKK (K) {20}
117: VRKKK KKKKKKKKKKKKKKKKKKK (K) {20}
118: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
119: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
120: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
121: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

122: KKKKK (K) {20} KKKAV
123: KKKKK (K) {20} KKKAVL

ABG92662 ck: 7676 len: 156 ! Abg92662 Human DNA-binding protein #8. 11/

(R,K) {20,20}
108: KLTWI KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
109: TTWIK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
110: TWIKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
111: WIKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
112: IKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
113: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
114: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
115: KKKKK (R,K) {20} KKKKKK GKGST

ABG92663 ck: 1736 len: 40 ! Abg92663 Human DNA-binding protein #9. 11/

(R,K) {20,20}
18: LPSLK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
19: PPSLK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
20: GSLKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
21: SLKKK KKKKKKKKKKKKKKKKKKKKKKKKK

ABG92665 ck: 1109 len: 98 ! Abg92665 Human DNA-binding protein #91. 11/

(R,K) {20,20}
53: QTNKT KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
54: TNNTK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
55: KNTKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
56: NTKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
57: TKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
58: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

1

(K) {20}
59: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
60: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
61: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
62: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
63: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
64: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
65: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
66: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
67: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
68: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20} GGRSR

ABG92667 ck: 8102 len: 111 ! Abg92667 Human DNA-binding protein #93. 11/

(R,K) {20,20}
78: EFHIL KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
79: FHILK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
80: HILKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
81: ILKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
82: LKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
83: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
84: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
85: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
86: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
87: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
88: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
89: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
90: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK

91: KKKK (K) {20} K
(K) {20}
92: KKKK KKKKKKKKKKKKKKKKK

ABG92668 ck: 8102 len: 111 ! Abg92668 Human DNA-binding protein #94. 11/

(R,K) {20,20}

(K) {20}

78: EFHIL KKKKKKKKKKKKKKKKK

(K) {20}

79: FHILK KKKKKKKKKKKKKKKKK

(K) {20}

80: HILK KKKKKKKKKKKKKKKKK

(K) {20}

81: ILKK KKKKKKKKKKKKKKKKK

(K) {20}

82: LKKK KKKKKKKKKKKKKKKKK

(K) {20}

83: KKKK KKKKKKKKKKKKKKKKK

(K) {20}

84: KKKK KKKKKKKKKKKKKKKKK

(K) {20}

85: KKKK KKKKKKKKKKKKKKKKK

(K) {20}

86: KKKK KKKKKKKKKKKKKKKKK

(K) {20}

87: KKKK KKKKKKKKKKKKKKKKK

(K) {20}

88: KKKK KKKKKKKKKKKKKKKKK

(K) {20}

89: KKKK KKKKKKKKKKKKKKKKK

(K) {20}

90: KKKK KKKKKKKKKKKKKKKKK

(K) {20}

91: KKKK KKKKKKKKKKKKKKKKK

(K) {20}

92: KKKK KKKKKKKKKKKKKKKKK

ABG92669 ck: 8319 len: 53 ! Abg92669 Human DNA-binding protein #95. 11/

(R,K) {20,20}

(K) {20}

13: RYKRP KKKKKKKKKKKKKKKKK

(K) {20}

14: YKPK KKKKKKKKKKKKKKKKK

(K) {20}

15: FKPK KKKKKKKKKKKKKKKKK

(K) {20}

16: KPKK KKKKKKKKKKKKKKKKK

(K) {20}

17: PKKK KKKKKKKKKKKKKKKKK

18: KKKK (K) {20} KKKK

19: KKKK (K) {20} KKKK

20: KKKK (K) {20} KKKK

21: KKKK (K) {20} KKKK

22: KKKK (K) {20} KKKK

23: KKKK (K) {20} KKKK

24: KKKK (K) {20} KKKK

25: KKKK (K) {20} KKKK

26: KKKK (K) {20} KKKK

27: KKKK (K) {20} KKKK

28: KKKK (K) {20} KKEG

29: KKKK (K) {20} KKEG

30: KKKK (K) {20} XEGX

ABG92671 ck: 7918 len: 80 ! Abg92671 Human DNA-binding protein #97. 11/

(R,K) {20,20}

(K) {20}

50: NVLTV KKKKKKKKKKKKKKKKK

51: VLTVK KKKKKKKKKKKKKKKKK

52: LTVK KKKKKKKKKKKKKKKKK

53: TVKK KKKKKKKKKKKKKKKKK

54: VKKK (K) {20} KKKK

55: KKKK (K) {20} KKKK

56: KKKK (K) {20} XXXX

ABG92673 ck: 4882 len: 41 ! Abg92673 Human DNA-binding protein #99. 11/

(R,K) {20,20}

(K) {20}

8: FYCFP KKKKKKKKKKKKKKKKK

(K) {20}

```
9: YCFEK KKKKKKKKKKKKKKKKKKKKK KKKK
    (K) {20}
10: CFFKK KKKKKKKKKKKKKKKKKKKKK KKKK
    (K) {20}
11: FPKKK KKKKKKKKKKKKKKKKKKKKK KKKK
    (K) {20}
12: FKKKK KKKKKKKKKKKKKKKKKKKKK KKKK
    (K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKK
    (K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKK
    (K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKK
    (K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKG
    (K) {20}
17: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKGX
    (K) {20}
18: KKKKK KKKKKKKKKKKKKKKKKKKKK KKGX
    (K) {20}
19: KKKKK KKKKKKKKKKKKKKKKKKKKK KKG
    (K) {20}
20: KKKKK KKKKKKKKKKKKKKKKKKKKK GX
    (K) {20}

ABG92674 ck: 5469 len: 63 1 Abg92674 Human DNA-binding protein #100. 11
(R, K) {20, 20}
(K) {20}
30: ITCLL KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
31: ICLLK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
32: CLKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
33: LKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
34: LKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
35: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
36: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
37: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
38: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
39: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKG
    (K) {20}
40: KKKKK KKKKKKKKKKKKKKKKKKKKK KKGX
    (K) {20}
```

```
41: KKKKK KKKKKKKKKKKKKKKKKKKKK KKG
    (K) {20}
42: KKKKK KKKKKKKKKKKKKKKKKKKKK XG
    (K) {20}

ABG92675 ck: 5075 len: 52 1 Abg92675 Human DNA-binding protein #101. 11
(R, K) {20, 20}
(K) {20}
30: FIVVX KKKKKKKKKKKKKKKKKKKKK KKK
    (K) {20}
31: IVVXX KKKKKKKKKKKKKKKKKKKKK KK
    (K) {20}
32: VVXXK KKKKKKKKKKKKKKKKKKKKK K
    (K) {20}
33: VXXKK KKKKKKKKKKKKKKKKKKKKK
    (K) {20}

ABG92676 ck: 5741 len: 47 1 Abg92676 Human DNA-binding protein #102. 11
(R, K) {20, 20}
(K) {20}
20: ILTTF KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
21: LTTFK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
22: TTFKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
23: TFKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
24: FKKKK KKKKKKKKKKKKKKKKKKKKK KKKK
    (K) {20}
25: KKKKK KKKKKKKKKKKKKKKKKKKKK KKK
    (K) {20}
26: KKKKK KKKKKKKKKKKKKKKKKKKKK KK
    (K) {20}
27: KKKKK KKKKKKKKKKKKKKKKKKKKK X
    (K) {20}

ABG92677 ck: 2868 len: 84 1 Abg92677 Human DNA-binding protein #103. 11
(R, K) {20, 20}
(K) {20}
53: KCTVB KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
54: CTYBK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
55: TYBKX KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
56: YBKXK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
57: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
58: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
59: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
    (K) {20}
```

(K) {20}
60: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
61: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}
62: KKKKK KKKKKKKKKKKKKKKKKKK KKK

ABG92678 ck: 4686 len: 73 1 Abg92678 Human DNA-binding protein #104. 11

(R,K) {20,20}

(K) {20}

41: YLKE KKKKKKKKKKKKKKKKKKK KKKK

42: LKKE KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

43: KKEK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

44: KEKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

45: EKKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

46: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

47: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

48: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

49: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

50: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

51: KKKK KKKKKKKKKKKKKKKKKKK KKK
(K) {20}

52: KKKK KKKKKKKKKKKKKKKKKKK KK
(K) {20}

53: KKKK KKKKKKKKKKKKKKKKKKK K
(K) {20}

54: KKKK KKKKKKKKKKKKKKKKKKK

ABG92679 ck: 6676 len: 74 1 Abg92679 Human DNA-binding protein #105. 11

(R,K) {20,20}

(K) {20}

47: LRTFQ KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

48: RTFQK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

49: TFOK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

50: FOKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

51: OKKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}
52: KKKK KKKKKKKKKKKKKKKKKKK KKK

ABG92680 ck: 2283 len: 54 1 Abg92680 Human DNA-binding protein #106. 11

(R,K) {20,20}

(K) {20}

32: IVFCF KKKKKKKKKKKKKKKKKKK KKK

33: VFCFK KKKKKKKKKKKKKKKKKKK KK
(K) {20}

34: PCFK KKKKKKKKKKKKKKKKKKK X

ABG92681 ck: 7503 len: 74 1 Abg92681 Human DNA-binding protein #107. 11

(R,K) {20,20}

(K) {20}

45: SHLTD KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

46: HLTDK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

47: LTDKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

48: TDKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

49: DKKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

50: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

51: KKKK KKKKKKKKKKKKKKKKKKK KKK
(K) {20}

52: KKKK KKKKKKKKKKKKKKKKKKK KKK
(K) {20}

53: KKKK KKKKKKKKKKKKKKKKKKK KK
(K) {20}

54: KKKK KKKKKKKKKKKKKKKKKKK K
(K) {20}

55: KKKK KKKKKKKKKKKKKKKKKKK

ABG92683 ck: 5199 len: 84 1 Abg92683 Human DNA-binding protein #109. 11

(R,K) {20,20}

(K) {20}

63: AMNAG KKKKKKKKKKKKKKKKKKK XG

ABG92684 ck: 7578 len: 31 1 Abg92684 Human DNA-binding protein #110. 11

(R,K) {20,20}

(K) {20}

6: LTELK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

7: TELK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}

8: ELEKK KKKKKKKKKKKKKKKKKKKKK KKKK
(K) {20}
9: LEKKK KKKKKKKKKKKKKKKKKKKKK KKK
(K) {20}
10: EKKKK KKKKKKKKKKKKKKKKKKKKK KK
(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKK X
(K) {20}

ABG92685 ck: 3915 len: 57 ! Abg92685 Human DNA-binding protein #11. 11
(R,K) {20,20}
31: KQLLK KKKKKKKKKKKKKKKKKKKKK KKKKG
(K) {20}
32: QLLLK KKKKKKKKKKKKKKKKKKKKK KKKGG
(K) {20}
33: LLLKK KKKKKKKKKKKKKKKKKKKKK KKGGF
(K) {20}
34: LLKKK KKKKKKKKKKKKKKKKKKKKK XGGF
(K) {20}

ABG92686 ck: 3679 len: 37 ! Abg92686 Human DNA-binding protein #12. 11
(R,K) {20,20}
15: ISPLT KKKKKKKKKKKKKKKKKKKKK KKK
(K) {20}
16: SPLTK KKKKKKKKKKKKKKKKKKKKK KK
(K) {20}
17: PLTKK KKKKKKKKKKKKKKKKKKKKK X
(K) {20}

ABG92687 ck: 657 len: 196 ! Abg92687 Human DNA-binding protein #13. 11
(R,K) {20,20}
169: FVXFE KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
170: VXFEE KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
171: XFEEK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
172: FEKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
173: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKK
(K) {20}
174: KKKKK KKKKKKKKKKKKKKKKKKKKK KKK
(K) {20}
175: KKKKK KKKKKKKKKKKKKKKKKKKKK KK
(K) {20}
176: KKKKK KKKKKKKKKKKKKKKKKKKKK X
(K) {20}

ABG92688 ck: 4672 len: 57 ! Abg92688 Human DNA-binding protein #14. 11
(R,K) {20,20}
(K) {20}

28: DKTFF KKKKKKKKKKKKKKKKKKKKK KKKK
(K) {20}
29: KTFHK KKKKKKKKKKKKKKKKKKKKK KKKKP
(K) {20}
30: TFFHK KKKKKKKKKKKKKKKKKKKKK KKKPG
(K) {20}
31: FHKKK KKKKKKKKKKKKKKKKKKKKK KKPGG
(K) {20}
32: HKKKK KKKKKKKKKKKKKKKKKKKKK XPGGG
(K) {20}

ABG92689 ck: 9656 len: 66 ! Abg92689 Human DNA-binding protein #15. 11
(R,K) {20,20}
38: MVISV KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
39: VISVK KKKKKKKKKKKKKKKKKKKKK KKKRE
(K) {20}
40: ISVKK KKKKKKKKKKKKKKKKKKKKK KKKRE
(K) {20}
41: SVKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
42: VKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
43: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(R,K) {20}

ABG92691 ck: 4665 len: 34 ! Abg92691 Human DNA-binding protein #17. 11
(R,K) {20,20}
10: PELLK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
11: ELLKK KKKKKKKKKKKKKKKKKKKKK KKKK
(K) {20}
12: LLLKK KKKKKKKKKKKKKKKKKKKKK KKK
(K) {20}
13: LLKKK KKKKKKKKKKKKKKKKKKKKK KK
(K) {20}
14: LKKKK KKKKKKKKKKKKKKKKKKKKK K
(K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKK
(K) {20}

ABG92692 ck: 7810 len: 64 ! Abg92692 Human DNA-binding protein #18. 11
(R,K) {20,20}
37: LKYFW KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
38: KYFWK KKKKKKKKKKKKKKKKKKKKK KKKKG
(K) {20}
39: YFWKK KKKKKKKKKKKKKKKKKKKKK KKKGX
(K) {20}
40: FWKKK KKKKKKKKKKKKKKKKKKKKK KKKGP
(K) {20}

41: WKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK XGXP

ABG92693 ck: 269 len: 76 ! Abg92693 Human DNA-binding protein #119. 11

(R, K) {20, 20}
 (K) {20}

37: TPSPRA KKKKKKKKKKKKKKKKKKKKK KKKKK

38: PSPRAK KKKKKKKKKKKKKKKKKKKKK KKKKK

39: SPRAKK KKKKKKKKKKKKKKKKKKKKK KKKKK

40: PAKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

41: AKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

42: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

43: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKI

44: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK XKKIK

ABG92694 ck: 8370 len: 45 ! Abg92694 Human DNA-binding protein #120. 11

(R, K) {20, 20}
 (K) {20}

17: APKTKQ KKKKKKKKKKKKKKKKKKKKK KKKKK

18: PKTKQK KKKKKKKKKKKKKKKKKKKKK KKKKK

19: KTKQKK KKKKKKKKKKKKKKKKKKKKK KKKKK

20: TKQKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

21: QKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

22: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

23: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

24: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

25: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

ABG92695 ck: 1663 len: 87 ! Abg92695 Human DNA-binding protein #121. 11

(R, K) {20, 20}
 (K) {20}

36: KWSXK KKKKKKKKKKKKKKKKKKKKK KKKKK

37: WSSXK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

38: SSXKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

39: SXKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

40: XKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

41: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

42: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

43: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

44: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

45: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

46: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

47: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

48: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

49: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

50: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

51: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

52: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

53: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

54: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

55: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

56: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

57: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

58: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKKK

59: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKRG

60: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKGX

61: KKKKK (K) {20}
 KKKKKKKKKKKKKKKKKKKKK KKKGP

62: KKKKK KKKKKKKKKKKKKKKKKKKKKK RQKPF
 (R,K) {20}
 63: KKKKK KKKKKKKKKKKKKKKKKKKKKK GKPFX

ABG92696 ck: 5607 len: 63 ! Abg92696 Human DNA-binding protein #122. 11

(R,K) {20,20}

(K) {20}

26: MVELE KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

27: VELEK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

28: ELEKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

29: LEKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

30: EKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

31: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

32: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

33: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

34: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

35: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

36: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

37: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

38: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

39: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

40: KKKKK KKKKKKKKKKKKKKKKKKKKKK GGPFF

ABG92697 ck: 5997 len: 58 ! Abg92697 Human DNA-binding protein #123. 11

(R,K) {20,20}

(K) {20}

28: RPTRP KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

29: PTPPK KKKKKKKKKKKKKKKKKKKKKK KKKKK

ABG92698 ck: 5764 len: 74 ! Abg92698 Human DNA-binding protein #124. 11

(R,K) {20,20}

(K) {20}

40: EPLSA KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

41: FLSAK KKKKKKKKKKKKKKKKKKKKKK KKKKK

42: LSARK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

43: SAKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

44: AKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

45: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

46: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

47: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

48: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

49: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

50: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

51: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

52: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

53: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

54: KKKKK KKKKKKKKKKKKKKKKKKKKKK X

ABG92699 ck: 7734 len: 97 ! Abg92699 Human DNA-binding protein #125. 11

(R,K) {20,20}

(K) {20}

57: RGCSY KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

58: GCSYK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

59: CSYKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

60: SYKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

61: YKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

62: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

63: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

64: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

65: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

66: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK

67: KKKKK (K) {20}
 68: KKKKK (K) {20}
 69: KKKKK (K) {20}
 70: KKKKK (K) {20}
 71: KKKKK (K) {20}
 72: KKKKK (K) {20}
 73: KKKKK (K) {20}
 74: KKKKK (K) {20}
 75: KKKKK (K) {20} XGX

ABG92700 ck: 3164 len: 181 1 Abg92700 Human DNA-binding protein #126. 11

154: TRKPE KKKKK (R,K) {20,20}
 155: RPEPE KKKKK (K) {20}
 156: KPEPE KKKKK (K) {20}
 157: PEPEK KKKKK (K) {20}
 158: EPEKK KKKKK (K) {20}
 159: KKKKK KKKKK (K) {20}
 160: KKKKK KKKKK (K) {20} X
 161: KKKKK KKKKK (K) {20} X

ABG92701 ck: 7117 len: 55 1 Abg92701 Human DNA-binding protein #127. 11

22: DDKKN KKKKK (R,K) {20,20}
 23: DKKNK KKKKK (K) {20}
 24: KKNKK KKKKK (K) {20}
 25: KNKKK KKKKK (K) {20} XGG
 (K) {20}

26: NKKKK KKKKK
 27: KKKKK (K) {20} XGGP

ABG92702 ck: 9316 len: 67 1 Abg92702 Human DNA-binding protein #128. 11

36: PTLQT KKKKK (R,K) {20,20}
 37: TLQTR KKKKK (R,K) {20}
 38: LQTRK KKKKK (K) {20}
 39: QTRKK KKKKK (K) {20}
 40: TRKKK KKKKK (K) {20} XGGP
 41: RKKKK KKKKK (K) {20} XGGP

ABG78915 ck: 8085 len: 154 1 Abg78915 Human breast tumour polypeptide #7

114: TQLPQ KKKKK (R,K) {20,20}
 115: QLPQK KKKKK (K) {20}
 116: LRPQK KKKKK (K) {20}
 117: RQPKK KKKKK (K) {20}
 118: QPKKK KKKKK (K) {20}
 119: KKKKK KKKKK (K) {20}
 120: KKKKK KKKKK (K) {20}
 121: KKKKK KKKKK (K) {20}
 122: KKKKK KKKKK (K) {20}
 123: KKKKK KKKKK (K) {20}
 124: KKKKK KKKKK (K) {20}
 125: KKKKK KKKKK (K) {20}
 126: KKKKK KKKKK (K) {20}
 127: KKKKK KKKKK (K) {20}

81: VLKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
82: LKKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
83: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
84: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
85: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
86: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}

1 ABG65251 ck: 8971 len: 108 ! Abg65251 Human albumin fusion protein #1926
(R, K) {20, 20}

78: TLLXL KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
79: LLLXL KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
80: LLLXL KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
81: XLKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
82: LKKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
83: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
84: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
85: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
86: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}

1 ABG65440 ck: 1431 len: 530 ! Abg65440 Human albumin fusion protein #2115
(R, K) {20, 20}

511: LHAP KKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}

1 ABG5896 ck: 5383 len: 86 ! Abg5896 Human peptide encoded by genome-de
(R, K) {20, 20}

15: RRRRG RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
16: RRRGR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
17: RRRGR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
18: RRRGR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}

19: GRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
20: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
21: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
22: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
23: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
24: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
25: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
26: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
27: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
28: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
29: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
30: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
31: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
32: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
33: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
34: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
35: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
36: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
37: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
38: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
39: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
40: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
41: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
42: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}
43: RRRRR RRRRRKKKKKKKKKKKKKKKKKK (R, K) {20}

44: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRRR RNRQT
45: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRRR NRQTK

ABG36760 ck: 1334 len: 86 ! Abg36760 Human peptide encoded by genome-de

57: EEEEG (R,K) {20,20} RRRRRRRRRRRRRRRRRRRR KKKKK
58: EEEGR (R,K) {20} RRRRRRRRRRRRRRRRRRRR KKKKK

59: EEEGR (R,K) {20} RRRRRRRRRRRRRRRRRRRR KKKKK
60: EGRRR (R,K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

61: GRRRK (R,K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
62: RRRKK (R,K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

63: RRRKK (R,K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
64: RRRKK (R,K) {20} KRRRRRRRRRRRRRRRRRRR KKK

65: KRRKK (R,K) {20} KRRRRRRRRRRRRRRRRRRR KK
66: KRRKK (R,K) {20} KRRRRRRRRRRRRRRRRRRR K

67: KRRKK (K) {20} KRRRRRRRRRRRRRRRRRRR

ABG36843 ck: 9082 len: 167 ! Abg36843 Human peptide encoded by genome-de

33: EEEGR (R,K) {20,20} RRRRRRRRRRRRRRRRRRRR RRGCG
34: EGRGR (R,K) {20} RRRRRRRRRRRRRRRRRRRR RGGGR

35: GRRGR (R,K) {20} RRRRRRRRRRRRRRRRRRRR GGGRR

ABG37280 ck: 2276 len: 89 ! Abg37280 Human peptide encoded by genome-de

23: EEEER (R,K) {20,20} KRRRRRRRRRRRRRRRRRRR EEEKK
46: KEEER (K) {20} KRRRRRRRRRRRRRRRRRRR KRKEE

47: KEERK (K) {20} KRRRRRRRRRRRRRRRRRRR KRKEE
48: EEEKK (R,K) {20} KRRRRRRRRRRRRRRRRRRR KEEEE

1

49: EEEKK (R,K) {20} KRRRRRRRRRRRRRRRRRRR EEEER

ABG37848 ck: 3607 len: 88 ! Abg37848 Human peptide encoded by genome-de

39: ERKRE (R,K) {20,20} KRRRRRRRRRRRRRRRRRRR KKKKK
40: RRRER (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

41: KREKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
42: REKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

43: EKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
44: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

45: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
46: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

47: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
48: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

49: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
50: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

51: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
52: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

53: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
54: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

55: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
56: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

57: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
58: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

59: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK
60: KKKKK (K) {20} KRRRRRRRRRRRRRRRRRRR KKKKK

61: KKKKK (K) {20}
62: KKKKK (K) {20}
63: KKKKK (K) {20}
64: KKKKK (K) {20}
65: KKKKK (K) {20}
66: KKKKK (K) {20}
67: KKKKK (K) {20}
68: KKKKK (K) {20}
69: KKKKK (K) {20}

ABG38450 ck: 3937 len: 85 1 Abg38450 Human peptide encoded by genome-de

1

1: (R,K) {20,20}
(K) {20}

2: K (K) {20}

3: KK (K) {20}

4: KKK (K) {20}

5: KKKK (K) {20}

6: KKKKK (K) {20}

7: KKKKK (K) {20}

8: KKKKK (K) {20}

9: KKKKK (K) {20}

10: KKKKK (K) {20}

11: KKKKK (K) {20}

ABG39191 ck: 2686 len: 71 1 Abg39191 Human peptide encoded by genome-de

1

(R,K) {20,20}

(K) {20}

(K) {20}

(K) {20}

21: KKKKK (K) {20}

22: KKKKK (K) {20}

23: KKKKK (K) {20}

24: KKKKK (K) {20}

25: KKKKK (R,K) {20}

26: KKKKK (R,K) {20}

27: KKKKK (R,K) {20}

28: KKKKK (R,K) {20}

29: KKKKK (R,K) {20}

30: KKKKK (R,K) {20}

31: KKKKK (R,K) {20}

32: KKKKK (R,K) {20}

33: KKKKK (R,K) {20}

34: KKKKK (R,K) {20}

35: KKKKK (R,K) {20}

36: KKKKK (R,K) {20}

37: KKKKK (R,K) {20}

38: KKKKK (R,K) {20}

39: KKKKK (R,K) {20}

40: KKKKK (R,K) {20}

41: KKKKK (R,K) {20}

42: KKKKK (R,K) {20}

43: KKKKK (R,K) {20}

44: KKKKK (R,K) {20}

45: KKKKK (R,K) {20}

46: KKKK (R,K){20} RKKKKKKKKKKKKKKKKKK KKKSA
47: KKKR (K){20} KKKKKKKKKKKKKKKKK KKSAA
48: KRKK (K){20} KKKKKKKKKKKKKKKKK KSAH
49: RKKK (K){20} KKKKKKKKKKKKKKKKK SAH
ABG40383 ck: 1560 len: 88 ! Abg40383 Human peptide encoded by genome-de
(R,K){20,20}
43: RRRG RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
44: ERGR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
45: RGRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
46: RGRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
47: GRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
48: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
49: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
50: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
51: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
52: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
53: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
54: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
55: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
56: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
57: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
58: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
59: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
60: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
61: RRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}

62: RRRR (R){20} RRRRRRRRRRRRRRRRRRR NTNN
ABG40676 ck: 2324 len: 36 ! Abg40676 Human peptide encoded by genome-de
(R,K){20,20}
16: KRRK (R,K){20} KRRRRRRRRRRRRRRRR R
17: ERKK (R,K){20} KRRRRRRRRRRRRRRRR
ABG43651 ck: 8343 len: 66 ! Abg43651 Human peptide encoded by genome-de
(R,K){20,20}
6: ETERE KKKRRKKKKKKKKKKKKKK KKKK
(R,K){20}
7: TERK KKKKKKKKKKKKKKKKKKK KKKK
(R,K){20}
8: EREK KKKKKKKKKKKKKKKKKKK KKKK
(R,K){20}
9: REKK KKKKKKKKKKKKKKKKKKK NKKK
ABG45640 ck: 2394 len: 57 ! Abg45640 Human peptide encoded by genome-de
(R,K){20,20}
20: EEEG RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
21: EEEG RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
22: EGRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
23: EGRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
24: GRRR RRRRRRRRRRRRRRRRRRR RRRR
(R){20}
25: RRRR RRRRRRRRRRRRRRRRRRR RRRR
ABG46635 ck: 3301 len: 52 ! Abg46635 Human peptide encoded by genome-de
(R,K){20,20}
12: KKKK (R,K){20} KKKKKKKKKKKKKKKKK KKKK
13: KKKK (R,K){20} KKKKKKKKKKKKKKKKK KKKK
14: KKKK (R,K){20} KKKKKKKKKKKKKKKKK KKKK
15: KKKK (R,K){20} KKKKKKKKKKKKKKKKK KKKK
16: NKKK (R,K){20} KKKKKKKKKKKKKKKKK KKKK
17: KKKK (R,K){20} KKKKKKKKKKKKKKKKK KKKK

18: KKKKK (R,K){20} KKKKK
19: KKKKK (R,K){20} KKKKK
20: KKKKK (R,K){20} KKKKK
21: KKKKK (R,K){20} KKKKK
22: KKKKK (R,K){20} KKKKK
23: KKKKK (R,K){20} KKKKK
24: KKKKK (R,K){20} KKKKK
25: KKKKK (R,K){20} KKKKK
26: KKKKK (R,K){20} KKKKK
27: KKKKK (R,K){20} KKKKK
28: KKKKK (R,K){20} KKKKK
29: KKKKK (R,K){20} KKKKK
30: KKKKK (R,K){20} KKKKK
31: KKKKK (R,K){20} KKKKK

ABG47062 ck: 4895 len: 51 ! Abg47062 Human peptide encoded by genome-de
(R,K){20,20}
(R,K){20}
18: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK
19: KKKKK (R,K){20} KKKKK
20: KKKKK (R,K){20} KKKKK
21: KKKKK (R,K){20} KKKKK
22: KKKKK (R,K){20} KKKKK
23: KKKKK (R,K){20} KKKKK
24: KKKKK (R,K){20} KKKKK
25: KKKKK (R,K){20} KKKKK
26: KKKKK (R,K){20} KKKKK
27: KKKKK (R,K){20} KKKKK
28: KKKKK (R,K){20} KKKKK
29: KKKKK (R,K){20} KKKKK
30: KKKKK (R,K){20} KKKKK
31: KKKKK (R,K){20} KKKKK

ABG7424 ck: 9633 len: 637 ! Abg7424 Human tumour marker protein se2-5.
(R,K){20,20}
(R,K){20}
618: GKKKK KKKKKKKKKKKKKKKKKKK KKKKK

AAE20631 ck: 8971 len: 108 ! Aae20631 Human gene 4 encoded secreted prot
(R,K){20,20}
(K){20}
78: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
79: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

```

80: LVLKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
81: XLKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
82: LKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
83: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
84: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
85: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
86: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

AAE20642 ck: 7046 len: 105 ! Aae20642 Human gene 4 encoded secreted prote
      (R, K) {20, 20}
      (K) {20}
78: TLVLV KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
79: LVLVK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
80: LVLKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
81: VLKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
82: LKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
83: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
84: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
85: KKKKK KKKKKKKKKKKKKKKKKKKKK K
      (K) {20}
86: KKKKK KKKKKKKKKKKKKKKKKKKKK

ABB9690 ck: 1158 len: 226 ! Abb9690 Human polypeptide SEQ ID NO 2066.
      (R, K) {20, 20}
      (K) {20}
198: GAESL KKKKKKKKKKKKKKKKKKKKK KGGNP
      (K) {20}
199: AESLK KKKKKKKKKKKKKKKKKKKKK KGRPX
      (K) {20}
200: ESLKK KKKKKKKKKKKKKKKKKKKKK GRPXK

AAE14544 ck: 7250 len: 35 ! Aae14544 Peptide p65 used in assay for detec
      (R, K) {20, 20}
      (K) {20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
2: K KKKKKKKKKKKKKKKKKKKKK KKKKK

```


3: KK (K) {20} KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
4: KK (K) {20} KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
5: KKKK (K) {20} KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
6: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
7: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
8: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
9: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
10: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
11: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
12: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
13: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK KKK
(K) {20}
14: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK KK
(K) {20}
15: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK K
(K) {20}
16: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK

AAU75162 ck: 5536 len: 84 ! AAU75162 Single-chain antigen-binding polyP
(R, K) {20, 20}
1: (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
2: K (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: KK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
4: KKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
5: KKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
6: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
7: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
8: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
9: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

10: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
11: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
12: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
13: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
14: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
15: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
16: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
17: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
18: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
19: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
20: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
21: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
22: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
23: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
24: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
25: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
26: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
27: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
28: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
29: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
30: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
31: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
32: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
33: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

6: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
7: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
8: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
9: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
10: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
11: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
12: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
13: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
14: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
15: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
16: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
17: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
18: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
19: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
20: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
21: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
22: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
23: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
24: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
25: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
26: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
27: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
28: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
29: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}

1
30: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
31: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
32: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
33: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
34: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
35: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
36: RKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
37: KRKKR KRKKRKRKKRKRKKR KRKKR (R,K) {20}
AAU75165 ck: 6708 len: 84 ! Aau75165 single-chain antigen-binding polyp
(R,K) {20,20}
1: RRRRRRRRRRRRRRRRRRR RRRR (R) {20}
2: R RRRRRRRRRRRRRRRRR RRRR (R) {20}
3: RR RRRRRRRRRRRRRRRRR RRRR (R) {20}
4: RRR RRRRRRRRRRRRRRRRR RRRR (R) {20}
5: RRR RRRRRRRRRRRRRRRRR RRRR (R) {20}
6: RRRR RRRRRRRRRRRRRRRRR RRRR (R) {20}
7: RRRR RRRRRRRRRRRRRRRRR RRRR (R) {20}
8: RRRR RRRRRRRRRRRRRRRRR RRRR (R) {20}
9: RRRR RRRRRRRRRRRRRRRRR RRRR (R) {20}
10: RRRR RRRRRRRRRRRRRRRRR RRRR (R) {20}
11: RRRR RRRRRRRRRRRRRRRRR RRRR (R) {20}
12: RRRR RRRRRRRRRRRRRRRRR RRRR (R) {20}
13: RRRR RRRRRRRRRRRRRRRRR RRRR (R) {20}
14: RRRR RRRRRRRRRRRRRRRRR RRRR (R) {20}
15: RRRR RRRRRRRRRRRRRRRRR RRRR (R) {20}

16: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
17: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
18: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
19: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
20: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
21: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
22: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
23: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
24: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
25: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
26: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
27: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
28: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
29: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
30: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
31: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
32: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
33: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
34: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
35: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
36: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
37: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}

AAE16634 ck: 1504 len: 32 ! Aae16634 Nuclear localisation signal peptid

2: K RKRKRRKRKRKRKRKRKRKR KRKR
 3: KR (R,K) {20} KRKRKRKRKRKRKRKRKR KRKR
 4: KRK RKRRKRKRKRKRKRKRKR KRKR (R,K) {20}
 5: KRKR KRKRKRKRKRKRKRKRKR KRKR (R,K) {20}
 6: KRKRK RKRRKRKRKRKRKRKRKR KRKR (R,K) {20}
 7: RKRKR KRKRKRKRKRKRKRKRKR KRKR (R,K) {20}
 8: KRKRK RKRRKRKRKRKRKRKRKR KRKR (R,K) {20}
 9: RKRKR KRKRKRKRKRKRKRKRKR KRKR (R,K) {20}
 10: KRKRK RKRRKRKRKRKRKRKRKR KRKR (R,K) {20}
 11: RKRKR KRKRKRKRKRKRKRKRKR KRKR (R,K) {20}
 12: KRKRK RKRRKRKRKRKRKRKRKR KRKR (R,K) {20}
 13: RKRKR KRKRKRKRKRKRKRKRKR KRKR (R,K) {20}

AAm48798 ck: 6221 len: 26 | Aam48798 Tumour-targeting peptide vector pE

1: (R,K) {20,20} (K) {20} RGDCF KKKKKKKKKKKKKKKKKKK

1: ABB44830 ck: 5275 len: 38 | Abb44830 Human protective sequence CNI-00745

1: (R,K) {20,20} (K) {20} MAQ KKKKKKKKKKKKKKKKKKK KKKKK

4: MAQ KKKKKKKKKKKKKKKKKKK KKKKK (K) {20}

5: MAQK KKKKKKKKKKKKKKKKKKK KKKKK (K) {20}

6: MAQK KKKKKKKKKKKKKKKKKKK KKKKK (K) {20}

7: AQKKK KKKKKKKKKKKKKKKKKKK KKKKK (K) {20}

8: QKKKK KKKKKKKKKKKKKKKKKKK KKKKK (K) {20}

9: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK (K) {20}

10: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK (K) {20}

11: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK (K) {20}

12: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK (K) {20}

13: KKKK (K) {20}
 14: KKKK (K) {20}
 15: KKKK (K) {20}
 16: KKKK (K) {20}
 17: KKKK (K) {20}

AAU69736 ck: 875 len: 46 1 Aau69736 Cell death protective sequence CNT

15: RKKK (R, K) {20, 20}
 16: EKKK (K) {20}
 17: KSKK (K) {20}
 18: KSKK (K) {20}
 19: SKKK (K) {20}
 20: KKKK (K) {20}
 21: KKKK (K) {20}
 22: KKKK (K) {20}
 23: KKKK (K) {20}
 24: KKKK (K) {20}
 25: KKKK (K) {20}

AAU69736 ck: 5862 len: 50 1 Aau69736 Cell death protective sequence CNT

11: ESKK (R, K) {20, 20}
 12: SALR (K) {20}
 13: ALGR (K) {20}
 14: LGRK (K) {20}
 15: GRKK (K) {20}
 16: RKKK (K) {20}

17: KKKK (K) {20}
 18: KKKK (K) {20}
 19: KKKK (K) {20}
 20: KKKK (K) {20}
 21: KKKK (K) {20}
 22: KKKK (K) {20}
 23: KKKK (K) {20}
 24: KKKK (K) {20}
 25: KKKK (K) {20}
 26: KKKK (K) {20}
 27: KKKK (K) {20}
 28: KKKK (K) {20}
 29: KKKK (K) {20}

ABG47914 ck: 5383 len: 86 1 Abg47914 Human liver peptide, SEQ ID No 265

15: RRRG (R, K) {20, 20}
 16: RRRG (R, K) {20}
 17: RRRG (R, K) {20}
 18: RRRR (R, K) {20}
 19: GRRR (R, K) {20}
 20: RRRR (R, K) {20}
 21: RRRR (R, K) {20}
 22: RRRK (R, K) {20}
 23: RRRK (R, K) {20}
 24: RKKK (R, K) {20}

[illegible]

59: EGRPR RRRKKKRRKKKKKKKKKKKKKKKKKK
 (R, K) {20}
60: EGRRR KKKKKRRKKKKKKKKKKKKKKKKKK
 (R, K) {20}
61: GRRRK KKKGRKKKKKKKKKKKKKKKKKK
 (R, K) {20}
62: RRRRK KKRKKKKKKKKKKKKKKKKKKKKKKKK
 (R, K) {20}
63: RRKKK KKRRKKKKKKKKKKKKKKKKKKKKKKKK
 (R, K) {20}
64: RKKKK RRRKKKKKKKKKKKKKKKKKKKKKKKK
 (R, K) {20}
65: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
 (R, K) {20}
66: KKKKK RRRKKKKKKKKKKKKKKKKKKKKKKKK
 (K) {20}
67: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

```

ABG48847 ck: 9082 len: 167 ! Abg
      (R,K){20,20}
33: EEGKG RRRRRRRRRRRRRRRRRR RRGGG
      (R,K){20}
34: EGRGR RRRRRRRRRRRRRRRRRR RGGGR
      (R,K){20}
35: GGRGR RRRRRRRRRRRRRRRRRR GGGGR
      (R,K){20}

```

```

ABG49354  ck: 2276  len: 89      !  Abg

      (R, K {20, 20})
23: EEEEE KKKKKKKRKRKKKKKKKK EEEKK
      (R, K {20})

46: KKEE KKKKKKKKKKKKKKKKKKK KRKEE
      (K {20})

47: KEEKK KKKKKKKKKKKKKKKKKKK RKEEE
      (K {20})

48: EEEKK KKKKKKKKKKKKKKKKKKK KEEEE
      (R, K {20})

49: EEEKK KKKKKKKKKKKKKKKKKKK EEEEE
      (R, K {20})

```

```
ABG45651 ck: 1939 len: 130 ! Abg
      (R,K){20,20}
42: EGKKE RRRRRRRRRRRRRRRRRRRRRRRRR
      (R,K){20}
43: GKKEK RRRRRRRRRRRRRRRRRRRRRRRRR
      (R,K){20}
44: KKKER RRRRRRRRRRRRRRRRRRRRRRRRR
      (R,K){20}
```

1

```
45: KEERRR RRRRRRRRRRRRRRRRRR RRRKK (R,K){20}
46: EERRRR RRRRRRRRRRRRRRRRRR RRRKK (R,K){20}
47: RRRRR RRRRRRRRRRRRRRRRRR RKKKK (R,K){20}
48: RRRRR RRRRRRRRRRRRRRRRRR RKKKK (R,K){20}
49: RRRRR RRRRRRRRRRRRRRRRRR RKKKK (R,K){20}
50: RRRRR RRRRRRRRRRRRRRRRRR RKKKE (R,K){20}
51: RRRRR RRRRRRRRRRRRRRRRRR RKKKE (R,K){20}
52: RRRRR RRRRRRRRRRRRRRRRRR RKKEE (R,K){20}
53: RRRRR RRRRRRRRRRRRRRRRRR RKEEE (R,K){20}
54: RRRRR RRRRRRRRRRRRRRRRRR EEEEE (R,K){20}

ABG4962 ck: 3607 len: 88 ! Abg4962 Human liver peptide, SEQ ID No 286
(R,K){20,20}
39: ERRRE RRRRRRRRRRRRRRRRRR RKKKK (R,K){20}
40: RKREK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
41: KRREK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
42: REKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
43: EKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
44: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
45: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
46: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
47: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
48: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
49: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
50: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
51: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
```

1

```
52: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
53: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
54: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
55: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
56: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
57: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
58: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
59: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
60: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
61: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
62: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
63: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
64: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
65: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
66: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
67: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
68: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}
69: KKKKK RRRRRRRRRRRRRRRRRR RKKKK (K){20}

ABG50530 ck: 3937 len: 85 ! Abg50530 Human liver peptide, SEQ ID No 291
(R,K){20,20}
1: KRRRRRRRRRRRRRRRRR RKKKK (R,K){20}
2: KRRRRRRRRRRRRRRRRR RKKKK (K){20}
3: KRRRRRRRRRRRRRRRRR RKKKK (K){20}
4: KRRRRRRRRRRRRRRRRR RKKKK (K){20}
5: KRRRRRRRRRRRRRRRRR RKKKK (K){20}
```

```

6: KKKKK (K) {20} KKKKK
7: KKKKK (K) {20} KKKKK
8: KKKKK (K) {20} KKKKK
9: KKKKK (K) {20} KKKKK
10: KKKKK (K) {20} KKKKK
11: KKKKK (K) {20} KKKKK
12: KKKKK (K) {20} KKKKK
13: KKKKK (K) {20} KKKKK
14: KKKKK (K) {20} KKKKK
15: KKKKK (K) {20} KKKKK
16: KKKKK (K) {20} KKKKK
17: KKKKK (K) {20} KKKKK
18: KKKKK (K) {20} KKKKK
19: KKKKK (K) {20} KKKKK
20: KKKKK (K) {20} KKKKK
21: KKKKK (K) {20} KKKKK
22: KKKKK (K) {20} KKKKK
23: KKKKK (K) {20} KKKKK
24: KKKKK (K) {20} KKKKK
25: KKKKK (K) {20} KKKKK
26: KKKKK (K) {20} KKKKK
27: KKKKK (K) {20} KKKKK
28: KKKKK (K) {20} KKKKK
29: KKKKK (K) {20} KKKKK
30: KKKKK (K) {20} KKKKK
31: KKKKK (K) {20} KKKKK
32: KKKKK (K) {20} KKKKK
33: KKKKK (K) {20} KKKKK
34: KKKKK (K) {20} KKKKK
35: KKKKK (K) {20} KKKKK
36: KKKKK (K) {20} KKKKK
37: KKKKK (K) {20} KKKKK
38: KKKKK (K) {20} KKKKK
39: KKKKK (K) {20} KKKKK
40: KKKKK (K) {20} KKKKK
41: KKKKK (K) {20} KKKKK
42: KKKKK (K) {20} KKKKK
43: KKKKK (K) {20} KKKKK
44: KKKKK (K) {20} KKKKK
45: KKKKK (K) {20} KKKKK
46: KKKKK (K) {20} KKKKK
47: KKKKK (K) {20} KKKKK
48: KKKKK (K) {20} KKKKK
49: KKKKK (K) {20} KKKKK
50: KKKKK (K) {20} KKKKK
51: KKKKK (K) {20} KKKKK
52: KKKKK (K) {20} KKKKK
53: KKKKK (K) {20} KKKKK
54: KKKKK (K) {20} KKKKK
55: KKKKK (K) {20} KKKKK
56: KKKKK (K) {20} KKKKK
57: KKKKK (K) {20} KKKKK
58: KKKKK (K) {20} KKKKK
59: KKKKK (K) {20} KKKKK

```

```

1
60: RRRR RRRRRRRRRRRRRRRR RRRTN
(R, K) {20}
61: RRRR RRRRRRRRRRRRRRRR RNTNN
(R, K) {20}
62: RRRR RRRRRRRRRRRRRRRR NTNNE
(R, K) {20}

ABG52597 ck: 2324 len: 36 ! Abg52597 Human liver peptide, SEQ ID No 312
(R, K) {20, 20}
16: KERKT KRRRRRRRRRRRRRRR R
(R, K) {20}
17: ERKTK KRRRRRRRRRRRRRRR

ABG55512 ck: 8343 len: 66 ! Abg55512 Human liver peptide, SEQ ID No 341
(R, K) {20, 20}
6: ETERE KKKRRKKKKKKKKKKKK KKKNK
(R, K) {20}
7: TEREK KKKRRKKKKKKKKKKKK KKKNK
(R, K) {20}
8: EREK KRRKKKKKKKKKKKKKK KKKKK
(R, K) {20}
9: REKKK KRRKKKKKKKKKKKKKK KKKKK

ABG58055 ck: 2394 len: 57 ! Abg58055 Human liver peptide, SEQ ID No 367
(R, K) {20, 20}
20: EEEEG RRRRRRRRRRRRRRRR RRRR
(R, K) {20}
21: EEGR RRRRRRRRRRRRRRRR RRRG
(R, K) {20}
22: EGRR RRRRRRRRRRRRRRRR RRRG
(R, K) {20}
23: EGRR RRRRRRRRRRRRRRRR RRRG
(R, K) {20}
24: GRRR RRRRRRRRRRRRRRRR RRRR
(R, K) {20}
25: RRRR RRRRRRRRRRRRRRRR RRRR

ABG58577 ck: 4228 len: 24 ! Abg58577 Human liver peptide, SEQ ID No 372
(R, K) {20, 20}
1: RRRRRRRRRRRRRRRR RRRR
(R, K) {20}
2: R RRRRRRRRRRRRRRRR RRRR
(R, K) {20}
3: RR RRRRRRRRRRRRRR RRRR
(R, K) {20}
4: RRR RRRRRRRRRRRRRR RRRR
(R, K) {20}

```


1
ABG59254 ck: 3301 len: 52 ! Abg59254 Human liver peptide, SEQ ID No 379
(R,K){20,20}
12: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
13: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
14: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
15: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
16: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
17: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
18: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
19: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
20: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
21: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
22: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
23: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
24: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
25: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
26: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
27: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
28: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
29: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
30: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
31: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
ABG59685 ck: 4895 len: 51 ! Abg59685 Human liver peptide, SEQ ID No 383
(R,K){20,20}
18: LFKPM KKKKKKKKKKKKKKKKKRR KKLTT
(R,K){20}
19: FKPMR KKKKKKKKKKKKKKKKKRR KLT TT

1
ABB96057 ck: 5626 len: 139 ! Abb96057 Human testicular antigen SEQ ID NC
(R,K){20,20}
111: IHLNL KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
112: HLNLK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
113: LNLKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
114: NLKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
1
ABB96575 ck: 4751 len: 80 ! Abb96575 Human testicular antigen SEQ ID NC
(R,K){20,20}
(K){20}
61: KKKFD KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
1
AAU87124 ck: 5599 len: 281 ! Aau87124 Novel central nervous system prote
(R,K){20,20}
(K){20}
248: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
249: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
250: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
251: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
252: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
253: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
254: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
255: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
256: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
257: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
258: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
259: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
260: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
261: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

262: KKKKK (K) {20}
KKKKKKKKKKKKKKKKKK

ABG00401 ck: 5116 len: 1,074 | Abg00401 Novel human diagnostic protein #39

(R, K) {20,20}

(R, K) {20}

609: RGSSS KKKKKKKKKKKKKKKKKRR KKKKK

(R, K) {20}

610: GSSSK KKKKKKKKKKKKKKKRRK NRKKK

ABG03974 ck: 5341 len: 99 | Abg03974 Novel human diagnostic protein #39

(R, K) {20,20}

(K) {20}

2: M KKKKKKKKKKKKKKKKKKK KKKKN

3: MK KKKKKKKKKKKKKKKKKKK KKKNS

4: MKK KKKKKKKKKKKKKKKKKKK KKNSS

5: MKKK KKKKKKKKKKKKKKKKKKK KNSOI

6: MKKKK KKKKKKKKKKKKKKKKKKK NSQID

ABG04391 ck: 9047 len: 139 | Abg04391 Novel human diagnostic protein #43

(R, K) {20,20}

(K) {20}

79: EEEEE KKKKKKKKKKKKKKKKKKK KKKKK

80: EEEEE KKKKKKKKKKKKKKKKKKK KKKKK

81: EEEEE KKKKKKKKKKKKKKKKKKK KKKKK

82: EEEEE KKKKKKKKKKKKKKKKKKK KKKKK

83: EEEEE KKKKKKKKKKKKKKKKKKK KKKKK

84: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

85: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

86: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

87: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

88: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

89: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

90: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

91: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

92: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

93: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

94: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

95: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

96: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

97: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

98: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

99: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

100: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

101: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

102: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

103: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

104: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

105: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

106: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

107: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

108: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

109: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

110: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

111: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

112: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

113: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

ABG05352 ck: 1276 len: 204 | Abg05352 Novel human diagnostic protein #53'

(R, K) {20,20}

(K) {20}

1

106: EEEEE KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 107: EEEEE KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 108: EEEEE KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 109: EEEEE KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 110: EEEEE KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 111: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 112: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 113: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20}
 114: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20}
 115: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20}
 116: KKKKK KKKKKKKKKKKKKKKKKKK KKEE
 (R, K) {20}
 117: KKKKK KKKKKKKKKKKKKKKKKKK KKEE
 (R, K) {20}
 118: KKKKK KKKKKKKKKKKKKKKKKKK EEEEE
 (R, K) {20}

ABG05367 ck: 6907 len: 808 1 Abg05367 Novel human diagnostic protein #53

1
 219: EEEEE RKKKKKKKKKKKKKKKKRR RRRR
 (R, K) {20}
 (R, K) {20}
 220: EEEEE RKKKKKKKKKKKKKKRRR RRRR
 (R, K) {20}
 221: EEEEE KKKKKKKKKKKKKKKRRR RRRR
 (R, K) {20}
 222: EEEEE KKKKKKKKKKKKKKKRRR RRRR
 (R, K) {20}
 223: EEEEE KKKKKKKKKKKKKKKRRR RRRR
 (R, K) {20}
 224: RRRR KKKKKKKKKKKKKKKRRR RRRR
 (R, K) {20}
 225: RRRR KKKKKKKKKKKKKKKRRR RRRR
 (R, K) {20}
 226: KKKKK KKKKKKKKKKKKKKKRRR KKKK
 (R, K) {20}
 227: KKKKK KKKKKKKKKKKKKKKRRR KKKK
 (R, K) {20}
 228: KKKKK KKKKKKKKKKKKKKKRRR KKKK
 (R, K) {20}

229: KKKKK RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 230: KKKKK RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 231: KKKKK RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 232: KKKKK RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 233: KKKKK RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 234: RRRR RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 235: RRRR RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 236: RRRR RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 237: RRRR RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 258: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 259: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 260: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}

ABG06375 ck: 7807 len: 2,570 1 Abg06375 Novel human diagnostic protein #63

1
 205: KKKSS RRRRRRRRRRRRRRRRRR RRRR
 (R, K) {20}
 (R, K) {20}
 206: KSSR RRRRRRRRRRRRRRRRRR RRRR
 (R, K) {20}
 207: KSSR RRRRRRRRRRRRRRRRRR RRRR
 (R, K) {20}
 208: SSRR RRRRRRRRRRRRRRRRRR RRRR
 (R, K) {20}
 ABG06513 ck: 2934 len: 154 1 Abg06513 Novel human diagnostic protein #65
 (R, K) {20, 20}
 (R, K) {20}
 49: KKKKT RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 50: KKKTR RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 51: KKKTR RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 52: KKKTR RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 53: KKKTR RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}
 54: KKKK RRRRRRRRRRRRRRRRRR KKKK
 (R, K) {20}

1

55: KKKKK (R,K){20} KKKKK
56: KRRKR (R,K){20} KKKKK
57: RKRRK (R,K){20} KKKKK
58: KRRKK (R,K){20} KKKKK
59: KRRKK (R,K){20} KKKKK
60: RKRRR (K){20} KKKKK
61: KKKRK (K){20} KKKKK
62: KRRKK (K){20} KKKKK
63: KRRKK (K){20} KKKKK
64: RKRRK (K){20} KKKKK
65: KKKKK (K){20} KKKKK

ABG07742 ck: 8672 len: 502 1 Abg07742 Novel human diagnostic protein #77

54: DDEEE (R,K){20,20} KKKKK
55: DEEER (R,K){20} KKKKK
56: EEEER (R,K){20} KKKKK
57: EERRR (R,K){20} KKKKK
58: ERRRR (R,K){20} KKKKK
59: RRRRR (R,K){20} KKKKK
60: RRRKK (R,K){20} KKKKK
61: RKRRK (R,K){20} KKKKK
62: KKKKK (R,K){20} KKKKK
63: KKKKK (R,K){20} KKKKK
64: KKKKK (R,K){20} KKKKK
65: KKKKK (R,K){20} KKKKK

1

66: KKKKK (R,K){20} KKKKK
67: KKKKK (R,K){20} KKKKK

ABG10052 ck: 7107 len: 39 1 Abg10052 Novel human diagnostic protein #10

2: Q (R,K){20,20} KKKKK
3: QK (K){20} KKKKK
4: QKK (K){20} KKKKK
5: QKKK (K){20} KKKKK
6: QKKKK (K){20} KKKKK
7: KKKKK (K){20} KKKKK

ABG10053 ck: 3274 len: 189 1 Abg10053 Novel human diagnostic protein #10

70: EKEKE (R,K){20,20} KKKKK
71: KEKEK (R,K){20} KKKKK
72: EKEKK (R,K){20} KKKKK
73: KEKKR (R,K){20} KKKKK
74: EKKRK (R,K){20} KKKKK
75: KKKRK (R,K){20} KKKKK
76: KKKKK (R,K){20} KKKKK
77: RKRRK (K){20} KKKKK
78: KKKRK (K){20} KKKKK
79: KKKKK (K){20} KKKKK

ABG11241 ck: 3870 len: 121 1 Abg11241 Novel human diagnostic protein #11

44: KKKKE (R,K){20,20} KKKKK
45: KKEEK (K){20} KKKKK

46: KEEKK KKKKKKKKKKKKKKKKKKKKK KKKKE
(K) {20}
47: EEEKK KKKKKKKKKKKKKKKKKKKKK KKKKE
(K) {20}
48: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKE
(K) {20}
49: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKE

ABG11242 ck: 4493 len: 100 ! Abg11242 Novel human diagnostic protein #11

1

(R,K) {20,20}
42: ETPSE KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
43: TPSEK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
44: PSEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
45: SEKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
46: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
47: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
48: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
49: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
50: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
51: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
52: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
53: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
54: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
55: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
56: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
57: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
58: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
59: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
60: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

1

61: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
62: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
63: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
64: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
65: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
66: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
67: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
68: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
69: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
70: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
71: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

ABG11245 ck: 2517 len: 85 ! Abg11245 Novel human diagnostic protein #11

(R,K) {20,20}
33: EEEEE KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
34: EEEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
35: EEEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
36: EEEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
37: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
38: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
39: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
40: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
41: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
42: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
43: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
44: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

45: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
46: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
47: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
48: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
49: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
50: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
51: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
52: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}

ABG11250 ck: 4343 len: 92 1 Abg11250 Novel human diagnostic protein #11

 (R,K) {20,20}
 (K) {20}
53: EKEKE KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
54: KEKEK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
55: EKEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
56: KEKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
57: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
58: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
59: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}

ABG11266 ck: 9563 len: 146 1 Abg11266 Novel human diagnostic protein #11

 (R,K) {20,20}
 (K) {20}
36: KEKKE KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
37: EKKEK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
38: KKEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
39: REKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
40: EKKKK KKKKKKKKKKKKKKKKKKKKK EEEEE
 (R,K) {20}
79: EEEEE KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
80: EEEKK KKKKKKKKKKKKKKKKKKKKK KKKKK

1

 (R,K) {20}
81: EEEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
82: EEEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
83: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
84: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
85: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
86: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
87: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
88: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
89: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
90: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
91: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
92: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
93: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}

ABG11277 ck: 8026 len: 1,080 1 Abg11277 Novel human diagnostic protein #11:

 (R,K) {20,20}
 (R,K) {20}
709: QKEKE KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
710: KEKEK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
711: EKEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
712: KEKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
713: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
714: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
715: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
716: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
717: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R,K) {20}
718: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

719: KKKK (R,K){20} KKKK
720: KKKK (R,K){20} KKKK
721: KKKK (R,K){20} KKKK
722: KKKK (R,K){20} KKKK
723: KKKK (R,K){20} KKKK
724: KKKK (R,K){20} KKKK
725: KKKK (R,K){20} KKKK
726: KKKK (R,K){20} KKKK
727: KKKK (R,K){20} EERR

1
ABG11732 ck: 2886 len: 56 i Abg11732 Novel human diagnostic protein #11
(R,K){20,20}
8: KRRG RRRRRRRRRRRRRRRRRR
(R){20}
9: RRRG RRRRRRRRRRRRRRRR
(R){20}
10: RRRG RRRRRRRRRRRRRRRR
(R){20}
11: RRRG RRRRRRRRRRRRRRRR
(R){20}
12: RRRG RRRRRRRRRRRRRRRR
(R){20}
13: RRRR RRRRRRRRRRRRRRRR

1
ABG11734 ck: 4548 len: 142 i Abg11734 Novel human diagnostic protein #11
(R,K){20,20}
109: QMLSV KKKKKKKKKKKKKKKK
(R,K){20}
110: MLSV KKKKKKKKKKKKKKKK
(R,K){20}
111: LSVK KKKKKKKKKKKKKKKK
(R,K){20}
112: SVKK KKKKKKKKKKKKKKKK
(R,K){20}
113: VKKK KKKKKKKKKKKKKKKK
(R,K){20}
114: KKKK KKKKKKKKKKKKKKKK
(R,K){20}
115: KKKK KKKKKKKKKKKKKKKK

116: KKKK (K){20} KKKK
117: KKKK (K){20} KKKK
118: KKKK (K){20} KKKK

1
ABG11738 ck: 3009 len: 567 i Abg11738 Novel human diagnostic protein #11
(R,K){20,20}
526: EEEE RRRRRRRRRRRRRRRR
(R,K){20}
527: EEEER RRRRRRRRRRRRRRRR
(R,K){20}
528: EEEER RRRRRRRRRRRRRRRR
(K){20}
529: EERR KKKKKKKKKKKKKKKK
(K){20}
530: EERR KKKKKKKKKKKKKKKK
(K){20}
531: RRRK KKKKKKKKKKKKKKKK
(K){20}
532: RRRK KKKKKKKKKKKKKKKK
(R,K){20}
533: KKKK KKKKKKKKKKKKKKKK
(R,K){20}
534: KKKK KKKKKKKKKKKKKKKK

1
ABG22512 ck: 8641 len: 856 i Abg22512 Novel human diagnostic protein #22
(R,K){20,20}
130: KKKF KKKKKKKKKKKKKKKK
(K){20}
131: KKKF KKKKKKKKKKKKKKKK
(K){20}
132: KKKF KKKKKKKKKKKKKKKK
(K){20}
133: KKKF KKKKKKKKKKKKKKKK
(K){20}
134: FKFK KKKKKKKKKKKKKKKK
(K){20}
135: KKKK KKKKKKKKKKKKKKKK
(K){20}
136: KKKK KKKKKKKKKKKKKKKK
(K){20}
1
ABG22638 ck: 4197 len: 896 i Abg22638 Novel human diagnostic protein #22
(R,K){20,20}
227: GREER RRRRRRRRRRRRRRRR

228: RERER RRRRRKRRRRKRRRRR RRRGX
 (R,K){20}
 229: EREER RRRKRRRRKRRRRRRR RRGXE
 (R,K){20}
 230: RERRR RRRKRRRRKRRRRRRR RGXEF
 (R,K){20}
 231: ERRRR RRRKRRRRKRRRRRRR GXEFL

1
 ABG26213 ck: 6773 len: 735 1 Abg26213 Novel human diagnostic protein #26
 (R,K){20,20}
 173: RGSSS KKKRRKKKKKKKKRRR KNRKK
 (R,K){20}
 174: GSSSK KKKRRKKKKKKKKRRK NRKKK
 (R,K){20}

1
 ABG26468 ck: 523 len: 124 1 Abg26468 Novel human diagnostic protein #26
 (R,K){20,20}
 91: EEEEE RRRRRKKKKKKKKKKK RKKKK
 (R,K){20}
 92: EEEEE RRRKKKKKKKKKKKKK KKKKK
 (R,K){20}
 93: EEEER RRRKKKKKKKKKKKKK KKKKR
 (R,K){20}
 94: EEEEE RRRKKKKKKKKKKKKK KKKRK
 (R,K){20}
 95: EEEEE RKKKKKKKKKKKKKKK KKKKK
 (R,K){20}
 96: RRRRR KKKKKKKKKKKKKKKK KKKKK
 (R,K){20}
 97: RRRRK KKKKKKKKKKKKKKKK RKKKK
 (R,K){20}
 98: RRRKK KKKKKKKKKKKKKKKK KKKKK
 (R,K){20}
 99: RRRKK KKKKKKKKKKKKKKKK KKKKK
 (R,K){20}
 100: RKKKK KKKKKKKKKKKKKKKK KKKKE
 (R,K){20}
 101: KKKKK KKKKKKKKKKKKKKKK KKKKE
 (R,K){20}
 102: KKKKK KKKKKKKKKKKKKKKK KKE
 (R,K){20}
 103: KKKKK KKKKKKKKKKKKKKKK KE
 (R,K){20}
 104: KKKKK KKKKKKKKKKKKKKKK E
 (R,K){20}

ABG26469 ck: 9923 len: 120 1 Abg26469 Novel human diagnostic protein #26
 (R,K){20,20}
 (R){20}

70: RRRRG RRRRRRRRRRRRRRRR RKKKK
 (R){20}
 71: RRRGR RRRRRRRRRRRRRRRR RKKKE
 (R){20}
 72: RRRGR RRRRRRRRRRRRRRRR KKKKE
 (R,K){20}
 73: RRRRR RRRRRRRRRRRRRR KKKKE
 (R,K){20}
 74: GRRRR RRRRRRRRRRRRRR KEEEE
 (R,K){20}
 75: RRRRR RRRRRRRRRRRRKKK EEEEE

1
 ABG26490 ck: 4146 len: 96 1 Abg26490 Novel human diagnostic protein #26
 (R,K){20,20}
 39: KKGEE RRRRRRRRRRRRRRRR RRRGR
 (R){20}
 40: KGEER RRRRRRRRRRRRRRRR RRRGR
 (R){20}
 41: GEERR RRRRRRRRRRRRRRR RGRGR
 (R){20}
 42: EERRR RRRRRRRRRRRRRRR RGRGR
 (R){20}
 73: GRGRG RRRRRRRRRRRRRRR RRRR
 (R){20}
 74: RGRGR RRRRRRRRRRRRRRR RRR
 (R){20}
 75: GRGRR RRRRRRRRRRRRRRR RR
 (R){20}
 76: RGRRR RRRRRRRRRRRRRRR R
 (R){20}
 77: GRRRR RRRRRRRRRRRRRRR

1
 ABG26491 ck: 8179 len: 109 1 Abg26491 Novel human diagnostic protein #26
 (R,K){20,20}
 75: RRRGE KKKRRKKKKKKKKKKK KKKKK
 (R,K){20}
 76: RRGEX KKKRRKKKKKKKKKKK KKKKK
 (R,K){20}
 77: RGEKK KRRKKKKKKKKKKKKK KKKKK
 (R,K){20}
 78: GEKKK KRRKKKKKKKKKKKKK KKKKK
 (R,K){20}
 79: EKKKK RKKKKKKKKKKKKKKK KKKKG
 (R,K){20}
 80: KKKKK KKKKKKKKKKKKKKKK KKKGN
 (K){20}
 81: KKKRK KKKKKKKKKKKKKKKK KKGNL
 (K){20}

1

82: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK KGNLS
 83: KKKKK (K) {20} KKKKKKKKKKKKKKKKKKK GNLST

ABG26492 ck: 5234 len: 68 ! Abg26492 Novel human diagnostic protein #26

(R,K) {20,20}

29: KKKEE (R) {20} RRRRRRRRRRRRRRRRRRR RRRRG
 30: KKEER (R) {20} RRRRRRRRRRRRRRRRRRR RRRGK
 31: KEERR (R) {20} RRRRRRRRRRRRRRRRRRR RRGKK
 32: EERRR (R) {20} RRRRRRRRRRRRRRRRRRR RGKGD
 33: ERRRR (R) {20} RRRRRRRRRRRRRRRRRRR GKKG

ABG26493 ck: 4204 len: 80 ! Abg26493 Novel human diagnostic protein #26

(R,K) {20,20}

27: EKEKE (R,K) {20} KRRRRRRRRRRRRRRRRRR RRRRR
 28: KEKEK (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 29: EKEKR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 30: KEKRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 31: EKRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 32: KRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 33: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 34: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 35: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 36: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 37: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 38: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 39: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 40: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 41: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR

1

42: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 43: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 44: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 45: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 46: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 47: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 48: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 49: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 50: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRT
 51: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRNTN
 52: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRNTN
 53: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR NTNNE

ABG26496 ck: 8829 len: 90 ! Abg26496 Novel human diagnostic protein #26

(R,K) {20,20}

61: EKKEE (R,K) {20} KRRRRRRRRRRRRRRRRRR RRRRR
 62: KKEEK (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 63: KEKRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 64: EEKRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 65: EKRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 66: KRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRRRR
 67: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRNN
 68: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RRN
 69: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR RN
 70: RRRRR (R) {20} RRRRRRRRRRRRRRRRRRR N

ABG26497 ck: 9186 len: 115 ! Abg26497 Novel human diagnostic protein #26

72: RRRR (R){20} RRRR
73: RRRR (R){20} RRRR
74: RRRR (R){20} RRRR
75: RRRR (R){20} RRRR
76: RRRR (R){20} RRRR
77: RRRR (R){20} RRRR
78: RRRR (R){20} RRRR
79: RRRR (R){20} RRRR
80: RRRR (R){20} RRRR
81: RRRR (R){20} RRRR
82: RRRR (R){20} RRRR
83: RRRR (R){20} RRRR
84: RRRR (R){20} RRRR
85: RRRR (R){20} RRRR
86: RRRR (R){20} TTTT

ABG26498 ck: 7156 len: 140 ! Abg26498 Novel human diagnostic protein #26

97: RGGG (R,K){20,20} RRRR
98: RGGG (R){20} RRRR
99: GGGG (R){20} RRRR
100: GGRG (R){20} RRRR
101: GRRR (R){20} RRRR
102: RRRR (R){20} RRRR
103: RRRR (R){20} RRRR
104: RRRR (R){20} RRRR

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105: RRRR RRRRRRRRRRRRRRRR RRRR      (R) {20}
106: RRRR RRRRRRRRRRRRRRRR RRRR      (R) {20}
107: RRRR RRRRRRRRRRRRRRRR RRRR      (R) {20}
108: RRRR RRRRRRRRRRRRRRRR RRRK      (R) {20}
109: RRRR RRRRRRRRRRRRRRRR RRKY      (R) {20}
110: RRRR RRRRRRRRRRRRRRRR RKYK      (R) {20}
111: RRRR RRRRRRRRRRRRRRRR RKYK      (R) {20}
112: RRRR RRRRRRRRRRRRRRRR KYKL      (R) {20}
113: RRRR RRRRRRRRRRRRRRK YKYLE      (R,K) {20}

ABG26500 ck: 7252 len: 78 ! ABg26500 Novel human diagnostic protein #26
(R,X) {20,20}
35: GEEE RRRRRRRRRRRRRRRR RRRR      (R) {20}
36: GEER RRRRRRRRRRRRRRRR RRRR      (R) {20}
37: EEEER RRRRRRRRRRRRRRRR RRRR      (R) {20}
38: EEER RRRRRRRRRRRRRRRR RRRR      (R) {20}
39: ERER RRRRRRRRRRRRRRRR RRRR      (R) {20}
40: RRRR RRRRRRRRRRRRRRRR RRRR      (R) {20}
41: RRRR RRRRRRRRRRRRRRRR RRRR      (R) {20}
42: RRRR RRRRRRRRRRRRRRRR RRRR      (R) {20}
43: RRRR RRRRRRRRRRRRRRRR RRRR      (R) {20}
44: RRRR RRRRRRRRRRRRRRRR RRRG      (R) {20}
45: RRRR RRRRRRRRRRRRRRRR RRRG      (R) {20}
46: RRRR RRRRRRRRRRRRRRRR RRGH      (R) {20}
47: RRRR RRRRRRRRRRRRRRRR RGHR      (R) {20}
48: RRRR RRRRRRRRRRRRRRRR GRHR      (R) {20}

```

1
ABG26501 ck: 1730 len: 182 ! Abg26501 Novel human diagnostic protein #26
(R,K){20,20}
72: KEEKE KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
73: EEEKE KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
74: EKEKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
75: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
76: EKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(R){20}
133: GRRRS RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
134: RRRSR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
135: RRSRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
136: RSRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
137: SRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
138: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
139: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
140: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
141: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
142: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
143: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
144: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
145: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
146: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
147: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
148: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
149: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
150: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}

1
ABG26502 ck: 3399 len: 101 ! Abg26502 Novel human diagnostic protein #26
(R,K){20,20}
59: EEEEG RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
60: EEEGR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
61: EEGRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
62: EGRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
63: GRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
64: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
65: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
66: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
67: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
68: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
69: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
1
ABG26505 ck: 4704 len: 93 ! Abg26505 Novel human diagnostic protein #26
(R,K){20,20}
60: KEEEG RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
61: EEEGR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
62: EEEGR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
63: EGGRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
64: GRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
65: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R){20}
66: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
67: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
68: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}
69: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRR
(R,K){20}

```

1
1
47: EKEKE KRRRRKKKKKKKKKKKKKKKKKK (R,K) {20}
48: KEKEK RRRRRKKKKKKKKKKKKKKKKKK (R,X) {20}
49: EKEKR KRRRRKKKKKKKKKKKKKKKKKK (R,K) {20}
50: KEKRR RRRRRKKKKKKKKKKKKKKKKKK (R,X) {20}
51: EKRRR KRRRRKKKKKKKKKKKKKKKKKK (R,K) {20}
52: KRRRK RRRRRKKKKKKKKKKKKKKKKKK (R,X) {20}
53: RRRRR KRRRRKKKKKKKKKKKKKKKKKK (K) {20}
54: KRRRK KRRRRKKKKKKKKKKKKKKKKKK (K) {20}
55: RRRRK KRRRRKKKKKKKKKKKKKKKKKK (K) {20}
56: KRRRK KRRRRKKKKKKKKKKKKKKKKKK (K) {20}
57: RRRRK KRRRRKKKKKKKKKKKKKKKKKK (K) {20}
58: KRRRK KRRRRKKKKKKKKKKKKKKKKKK (R,K) {20}
59: KRRRK KRRRRKKKKKKKKKKKKKKKKKK (R,K) {20}
60: KRRRK KRRRRKKKKKKKKKKKKKKKKKK (R,K) {20}
61: KRRRK KRRRRKKKKKKKKKKKKKKKKKK (R,K) {20}
62: KRRRK KRRRRKKKKKKKKKKKKKKKKKK (R,K) {20}

ABG26507 ck: 9838 len: 109 1 Abg26507 Novel human diagnostic protein #26
1
60: EEEEE RRRRRRRRRRRRRRRRRRRRRRR (R,K) {20,20}
61: EEEER RRRRRRRRRRRRRRRRRRRRRRR (R) {20}
62: EEEER RRRRRRRRRRRRRRRRRRRRRRR (R) {20}
63: EEEER RRRRRRRRRRRRRRRRRRRRRRR (R) {20}
64: EEEER RRRRRRRRRRRRRRRRRRRRRRR (R,K) {20}
65: EEEER RRRRRRRRRRRRRRRRRRRRRRR (R,K) {20}

```

1

ABG26508 ck: 4488 len: 121 ! Abg26508 Novel human diagnostic protein #26

56: DDEE RRRKKKKKKKKKKKKKK (R,K) {20,20}
(R,K) {20}

57: DEER RRRKKKKKKKKKKKKKK (R,K) {20}

58: EEER RKKKKKKKKKKKKKKKK (R,K) {20}

59: EERR RKKKKKKKKKKKKKKKK (R,K) {20}

60: EERR RKKKKKKKKKKKKKKKK (R,K) {20}

61: RRRK RKKKKKKKKKKKKKKKK (R,K) {20}

62: RRRK RKKKKKKKKKKKKKKKK (R,K) {20}

63: RRRK RKKKKKKKKKKKKKKKK (R,K) {20}

64: RRRK RKKKKKKKKKKKKKKKK (R,K) {20}

65: RRRK RKKKKKKKKKKKKKKKK (R,K) {20}

66: RRRK RKKKKKKKKKKKKKKKK (R,K) {20}

67: RRRK RKKKKKKKKKKKKKKKK (R,K) {20}

68: RRRK RKKKKKKKKKKKKKKKK (R,K) {20}

69: RRRK RKKKKKKKKKKKKKKKK (R,K) {20}

1

ABG26510 ck: 3426 len: 74 ! Abg26510 Novel human diagnostic protein #26

47: RRRS RRRRRRRRRRRRRRRR (R,K) {20,20}
(R) {20}

48: RRRS RRRRRRRRRRRRRRRR (R) {20}

49: RRRS RRRRRRRRRRRRRRRR (R) {20}

50: RRRS RRRRRRRRRRRRRRRR (R) {20}

51: RRRS RRRRRRRRRRRRRRRR (R) {20}

1

ABG26513 ck: 3117 len: 265 ! Abg26513 Novel human diagnostic protein #26

199: EEEE RRRRRRRRRRRRRRRR (R,K) {20,20}
(R,K) {20}

222: KKEE RRRRRRRRRRRRRRRR (R,K) {20}

223: KEEBK KKKKKKKKKKKKKKKKK RKEEB
(R,K){20}
224: EEEKK KKKKKKKKKKKKKKKKK KEEEB
(R,K){20}
225: EEEKK KKKKKKKKKKKKKKKKK EEEEE

ABG26514 ck: 9106 len: 218 | Abg26514 Novel human diagnostic protein #26

(R,K){20,20}
182: EEEBK RRRRRRRRRRRRRRRRR RRRR
(R){20}
183: EERGR RRRRRRRRRRRRRRRRR RRRR
(R){20}
184: RRGRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
185: RGRRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
186: GRRRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
187: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
188: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
189: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
190: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
191: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R,K){20}
192: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R,K){20}
193: RRRRR RRRRRRRRRRRRRRRRR RRRR

ABG26515 ck: 7288 len: 389 | Abg26515 Novel human diagnostic protein #26

(R,K){20,20}
259: KKEES RKKKKKKKKKKKKKKKK KKKR
(R,K){20}
260: KEESR KKKKKKKKKKKKKKKKK KKKR
(K){20}
261: EESRK KKKKKKKKKKKKKKKKK KRRN
(K){20}
262: ESRKK KKKKKKKKKKKKKKKKK RKNK
(R,K){20}
263: SRKKK KKKKKKKKKKKKKKKKK KRNK
(R,K){20}
264: RKKKK KKKKKKKKKKKKKKKKK RNKK
(R,K){20}

265: KKKKK KKKKKKKKKKKKKKKKK NKKK

ABG26516 ck: 2295 len: 91 | Abg26516 Novel human diagnostic protein #26

(R,K){20,20}
45: EEEEE RRRRRRRRRRRRRRRRR RRRR
(R,K){20}
46: EEEER RRRRRRRRRRRRRRRRR RRRR
(R,K){20}
47: EEEER RRRRRRRRRRRRRRRRR RRRR
(R,K){20}
48: EEEEE RRRRRRRRRRRRRRRRR RRRR
(R,K){20}
49: EEEEE RRRRRRRRRRRRRRRRR RRRR
(R,K){20}
50: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R,K){20}
51: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R,K){20}
52: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R,K){20}
53: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R,K){20}
54: RRRRR RRRRRRRRRRRRRRRRR RRRR

ABG26518 ck: 8431 len: 761 | Abg26518 Novel human diagnostic protein #26

(R,K){20,20}
725: EEEEE RRRRRRRRRRRRRRRRR RRRR
(R){20}
726: EEEEE RRRRRRRRRRRRRRRRR RRRR
(R){20}
727: EEEEE RRRRRRRRRRRRRRRRR RRRR
(R){20}
728: EEEEE RRRRRRRRRRRRRRRRR RRRR
(R){20}
729: EEEEE RRRRRRRRRRRRRRRRR RRRR
(R){20}
730: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
731: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
732: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
733: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
734: RRRRR RRRRRRRRRRRRRRRRR RRRR
(R){20}
735: RRRRR RRRRRRRRRRRRRRRRR RRRR

76: RRRRR RRRRRRRRRRRRRRRRRR RKKI
(R) {20}
77: RRRRR RRRRRRRRRRRRRRRRRR RKI
(R) {20}
78: RRRRR RRRRRRRRRRRRRRRRRR KI
(R,K) {20}
79: RRRRR RRRRRRRRRRRRRRRRRR I

ABg26521 ck: 482 len: 367 ! Abg26521 Novel human diagnostic protein #26

1

(R,K) {20,20}
317: EEEEG KKKKKKKKKKKKKKKKKK KKKK
(K) {20}
318: EEEGK KKKKKKKKKKKKKKKKK KKKR
(K) {20}
319: EEEGK KKKKKKKKKKKKKKKKK KKKR
(K) {20}
320: EGGKK KKKKKKKKKKKKKKKKK KRRK
(K) {20}
321: GKKKK KKKKKKKKKKKKKKKKK RRRK
(K) {20}
322: KKKKK KKKKKKKKKKKKKKKKK RRRK
(R,K) {20}
323: KKKKK KKKKKKKKKKKKKKKKK RKKK
(R,K) {20}
324: KKKKK KKKKKKKKKKKKKKKRR KKKK
(R,K) {20}
325: KKKKK KKKKKKKKKKKKKKKRR KKEQ
(R,K) {20}
326: KKKKK KKKKKKKKKKKKKRRKK KKEQ
(R,K) {20}
327: KKKKK KKKKKKKKKKKKKRRKK KEGQL
(R,K) {20}
328: KKKKK KKKKKKKKKKKRRKKK EGGLE
(R,K) {20}

ABg26522 ck: 2060 len: 152 ! Abg26522 Novel human diagnostic protein #26

1

(R,K) {20,20}
46: EEEEE RRRRRRRRRRRRRRRRRR RKKE
(R) {20}
47: EEEER RRRRRRRRRRRRRRRRRR RKKEG
(R) {20}
48: EEEER RRRRRRRRRRRRRRRRRR KKEGE
(R,K) {20}
49: EEEER RRRRRRRRRRRRRRRRRR KEGEE
(R,K) {20}
50: ERRRR RRRRRRRRRRRRRRRRRR EGEEG
(R) {20}
96: KKEEG RRRRRRRRRRRRRRRRRR RRRRR

(R) {20}
97: KEEGR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
98: EEEGR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
99: EGGRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
100: GRRRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
101: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
102: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
103: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
104: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
105: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
106: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
107: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
108: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
109: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
110: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R) {20}
111: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R,K) {20}
112: RRRRR RRRRRRRRRRRRRRRRRR RRRR

(R,K) {20}
113: RRRRR RRRRRRRRRRRRRRRRRR RRRR

ABg26525 ck: 5433 len: 103 ! Abg26525 Novel human diagnostic protein #26

1

(R,K) {20,20}
3: RD RRRKKKKKKKKKKKKKKKK EEEEE

1

ABg26526 ck: 1887 len: 115 ! Abg26526 Novel human diagnostic protein #26
(R,K) {20,20}
48: EEEEE RRRRRRRRRRRRRRRRRR KRRK
(R) {20}
49: EEEER RRRRRRRRRRRRRRRRRR KREKK
(R,K) {20}
50: EEEER RRRRRRRRRRRRRRRRRR REKKK
(R,K) {20}
51: EEEER RRRRRRRRRRRRRRRRRR EKKS

1
ABG26527 ck: 5586 len: 122 ! Abg26527 Novel human diagnostic protein #26
(R,K){20,20}
(R,K){20}

82: EEEEE KKKKKKKKKKKKKRRRRRR GRRRM

1
ABG26528 ck: 2237 len: 215 ! Abg26528 Novel human diagnostic protein #26
(R,K){20,20}
(R,K){20}

87: EKEKE KKKKKKKKKKKKKKKKKKK KKKKK

(R,K){20}

88: KEKEK KKKKKKKKKKKKKKKKKKK KKKKK

(R,K){20}

89: EKEKK RKKKKKKKKKKKKKKKKKK KKKKK

(R,K){20}

90: KEKKR KKKKKKKKKKKKKKKKKKK KKKKK

(R,K){20}

91: EKRRK KKKKKKKKKKKKKKKKKKK KKKKK

(R,K){20}

92: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(R,K){20}

93: KKKKK RKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

94: RKKRR KKKKKKKKKKKKKKKKKKK KKEEE

(K){20}

95: KKKKK KKKKKKKKKKKKKKKKKKK KEEEE

(K){20}

96: KKKKK KKKKKKKKKKKKKKKKKKK EEEEE

(R){20}

167: GRRRG RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

168: RRRGR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

169: RRRGR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

170: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

171: GRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

172: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

173: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

174: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

175: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

176: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

1
177: RRRRR RRRRRRRRRRRRRRRRRRR RRKRR
ABG26530 ck: 5729 len: 404 ! Abg26530 Novel human diagnostic protein #26
(R,K){20,20}
(R){20}

366: EEEEE RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

367: EEEER RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

368: EEEER RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

369: EEEER RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

370: ERRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

371: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

372: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

373: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

374: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

375: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

376: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

377: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

378: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R,K){20}

379: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

1
ABG26531 ck: 7434 len: 126 ! Abg26531 Novel human diagnostic protein #26
(R,K){20,20}
(R){20}

48: EEKDE RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

49: EKDER RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

50: KDERR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

51: DEERR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

52: EERRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

53: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

(R){20}

54: RRRRR RRRRRRRRRRRRRRRRRRR RRRRR

1

55: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

56: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

57: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

ABG26532 ck: 9650 len: 225 ! Abg26532 Novel human diagnostic protein #26

152: RRRRR (R,K){20,20} RRRRRRRRRRRRRRRRRRR RRRRR

153: RRRRR (R,K){20} RRRRRRRRRRRRRRRRRRR RRRRR

154: RRRRR (R,K){20} RRRRRRRRRRRRRRRRRRR RRRRR

155: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

156: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

157: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

158: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

159: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

160: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

161: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

162: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

163: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

164: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

165: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

166: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

167: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

ABG26533 ck: 5900 len: 101 ! Abg26533 Novel human diagnostic protein #26

51: RRRRR (R,K){20,20} RRRRRRRRRRRRRRRRRRR RRRRR

52: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

53: RRRRR (R){20} RRRRRRRRRRRRRRRRRRR RRRRR

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54: RTRRR RRRRRRRRRRRRRRRRRR RRKKK      (R) {20}
55: TRRRR RRRRRRRRRRRRRRRRRR RKKKP      (R) {20}
56: RRRRR RRRRRRRRRRRRRRRRRR KKCP      (R) {20}
57: RRRRR RRRRRRRRRRRRRRRRRR KKPTS      (R,K) {20}
58: RRRRR RRRRRRRRRRRRRRRR KPTSR      (R,K) {20}
59: RRRRR RRRRRRRRRRRRRRRR RPSRV      (R,K) {20}

Abg26534 ck: 1919 len: 113 1 Abg26534 Novel human diagnostic protein #26

1
(R,K) {20,20}
31: KKKKE KKKKKRRRRRRRRRRRRR KKKKK      (R,K) {20}
32: KKKKK KKKKKRRRRRRRRRRRRR KKKKK      (R,K) {20}
33: KKKKK KKKRRRRRRRRRRRRRRR KKKKK      (R,K) {20}
34: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (R,K) {20}
35: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (R,K) {20}
36: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (R,K) {20}
37: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (R,K) {20}
38: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (R,K) {20}
39: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (R,K) {20}
40: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (K) {20}
41: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (K) {20}
42: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (K) {20}
43: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (K) {20}
44: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (K) {20}
45: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (K) {20}
46: KKKRR KKKRRRRRRRRRRRRRRR KKKKK      (K) {20}
47: KKKRR KKKRRRRRRRRRRRRRRR KKKRR      (K) {20}

```

48: KKKK (K){20} KKKK
49: KKKK (K){20} KKKK
50: KKKK (R,K){20} KKKK
51: KKKK (R,K){20} KKKK
52: KKKK (R,K){20} KKKK
53: KKKK (R,K){20} KKKK
54: KKKK (R,K){20} KKKK
55: KKKK (R,K){20} KKKK
ABG26535 ck: 2214 len: 472 ! Abg26535 Novel human diagnostic protein #26
(R,K){20,20}
(R){20}

1

415: EEEE (R,K){20,20} RRRR
416: EEEE (R){20} RRRR
417: EEEE (R){20} RRRR
418: EEEE (R){20} RRRR
419: EEEE (R){20} RRRR
420: EEEE (R){20} RRRR
421: EEEE (R){20} RRRR
422: EEEE (R){20} RRRR
423: EEEE (R){20} RRRR
424: EEEE (R){20} RRRR
425: EEEE (R){20} RRRR
426: EEEE (R){20} RRRR
427: EEEE (R){20} RRRR
428: EEEE (R){20} RRRR
429: EEEE (R){20} RRRR

430: RRRR (R){20} RRRR
431: RRRR (R){20} RRRR
432: RRRR (R){20} RRRR
433: RRRR (R){20} RRRR
434: RRRR (R){20} RRRR
435: RRRR (R){20} RRRR
436: RRRR (R){20} RRRR
437: RRRR (R){20} RRRR
438: RRRR (R){20} RRRR
ABG26537 ck: 2429 len: 573 ! Abg26537 Novel human diagnostic protein #26
(R,K){20,20}
(R){20}

1

405: GAGG (R,K){20,20} RRRR
406: AEGG (R){20} RRRR
407: EGGG (R){20} RRRR
408: EGGG (R,K){20} RRRR
409: EGGG (R,K){20} RRRR
410: EGGG (R,K){20} RRRR
411: EGGG (R,K){20} RRRR
412: EGGG (R,K){20} RRRR
413: EGGG (R,K){20} RRRR
414: EGGG (R,K){20} RRRR
415: EGGG (R,K){20} RRRR
416: EGGG (R,K){20} RRRR
417: EGGG (R,K){20} RRRR
418: EGGG (R,K){20} RRRR
419: EGGG (R,K){20} RRRR
420: EGGG (R,K){20} RRRR
421: EGGG (R,K){20} RRRR
422: EGGG (R,K){20} RRRR
423: EGGG (R,K){20} RRRR
424: EGGG (R,K){20} RRRR
425: EGGG (R,K){20} RRRR
426: EGGG (R,K){20} RRRR
427: EGGG (R,K){20} RRRR
428: EGGG (R,K){20} RRRR
429: EGGG (R,K){20} RRRR
ABG26538 ck: 5732 len: 98 ! Abg26538 Novel human diagnostic protein #26
(R,K){20,20}
(R,K){20}

1

```

61: EKETK RRRRRKKKKKKKKKKKKKK KKKK
      (R, X) {20}
62: KETKR RRRRRKKKKKKKKKKKKKK KKKK
      (R, X) {20}
63: ETKR RRRKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
64: TKRR RRRKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
65: KRRR RKKKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
66: RRRR KKKKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
67: RRRR KKKKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
68: RRRK KKKRKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
69: RRRK KKRKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
70: RKKK KRRKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
71: KKKK RKKKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
72: KKKR KKKKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
73: KKKR KKKKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
74: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
75: KRRK KKKKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
76: RKKK KKKKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}
77: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (R, X) {20}

ABG26539 ck: 3545 len: 89 ; Abg26539 Novel human diagnostic protein #26
      (R, X) {20, 20}
28: EEEE RRRRRRRRRRRRRRRRRRR RRRR
      (R) {20}
29: EEEER RRRRRRRRRRRRRRRRRRR RRRR
      (R) {20}
30: EEER RRRRRRRRRRRRRRRRRRR RRRR
      (R) {20}
31: EERR RRRRRRRRRRRRRRRRRRR RRRR
      (R) {20}
32: ERRR RRRRRRRRRRRRRRRRRRR RRRR
      (R) {20}
33: RRRR RRRRRRRRRRRRRRRRRRR RRRR
      (R) {20}

```

[illegible]

```

265: RRRWR RRRRRRXXXXXXXXXXXXX KKKK      (R,K) {20}
266: RRWR RRRRRRXXXXXXXXXXXXX KKKK      (R,K) {20}
267: RWRR RRRRRRXXXXXXXXXXXXX KKKK      (R,K) {20}
268: WRRR RRRRRRXXXXXXXXXXXXX KKKK      (R,K) {20}
269: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (R,K) {20}
270: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (R,K) {20}
271: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (R,K) {20}
272: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (R,K) {20}
273: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (K) {20}
274: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (K) {20}
275: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (K) {20}
276: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (K) {20}
277: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (K) {20}
278: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (K) {20}
279: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (K) {20}
280: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (K) {20}
281: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (K) {20}
282: RRRR RRRRRRXXXXXXXXXXXXX KKKK      (K) {20}
ABG6545 ck: 135 len: 119 ! Abg26545 Novel human diagnostic protein #26
          (R,K) {20,20}
88: EKEE KKKKXXXXXXXXXXXXX KKKK      (K) {20}
89: KEKE KKKKXXXXXXXXXXXXX KKKK      (K) {20}
90: EEKE KKKKXXXXXXXXXXXXX KKKK      (K) {20}
91: KEKE KKKKXXXXXXXXXXXXX KKKK      (K) {20}
92: EKKE KKKKXXXXXXXXXXXXX KKKK      (K) {20}

```

[illegible]

18: KKKKK (R,K) {20} KKKKK
 19: KKKKK (R,K) {20} KKKKK
 20: KKKKK (R,K) {20} KKKKK
 21: KKKKK (R,K) {20} KKKKK

ABG26717 ck: 4415 len: 78 | Abg26717 Novel human diagnostic protein #26

37: OMLSV (R,K) {20,20} KKKKK
 38: MLSVK (R,K) {20} KKKKK
 39: LSVKK (R,K) {20} KKKKK
 40: SVKKK (R,K) {20} KKKKK
 41: VKKKK (R,K) {20} KKKKK
 42: KKKKK (R,K) {20} KKKKK
 43: KKKKK (R,K) {20} KKKKK
 44: KKKKK (R,K) {20} KKKKK
 45: KKKKK (R,K) {20} KKKKK
 46: KKKKK (R,K) {20} KKKKK
 47: RKKKK (R,K) {20} KKKKK
 48: KKKKK (R,K) {20} KKKKK
 49: KKKKK (R,K) {20} KKKKK
 50: KKKKK (R,K) {20} KKKKK
 51: KKKKK (R,K) {20} KKKKK
 52: KKKKK (R,K) {20} KKKKK
 53: KKKKK (R,K) {20} KKKKK
 54: KKKKK (R,K) {20} KKKKK

ABG26718 ck: 9531 len: 141 | Abg26718 Novel human diagnostic protein #26

80: EVARP (R,K) {20,20} KKKKK
 81: VAPRR (R,K) {20} KKKKK
 82: APRRK (R,K) {20} KKKKK
 83: RPRKK (R,K) {20} KKKKK
 84: PRKKK (R,K) {20} KKKKK
 85: RKKKK (R,K) {20} KKKKK
 86: KKKKK (R,K) {20} KKKKK
 87: KKKKK (R,K) {20} KKKKK
 88: KKKKK (R,K) {20} KKKKK
 89: KKKKK (R,K) {20} KKKKK

ABG26719 ck: 72 len: 83 | Abg26719 Novel human diagnostic protein #26

42: ETPSE (R,K) {20,20} KKKKK
 43: TPSEK (R,K) {20} KKKKK
 44: PSEKK (R,K) {20} KKKKK
 45: SEKKK (R,K) {20} KKKKK
 46: EKKKK (R,K) {20} KKKKK
 47: KKKKK (R,K) {20} KKKKK
 48: KKKKK (R,K) {20} KKKKK
 49: KKKKK (R,K) {20} KKKKK
 50: KKKKK (R,K) {20} KKKKK
 51: KKKKK (R,K) {20} KKKKK
 52: KKKKK (R,K) {20} KKKKK
 53: KKKKK (R,K) {20} KKKKK
 54: KKKKK (R,K) {20} KKKKK

1

ABG26720 ck: 3781 len: 57 i Abg26720 Novel human diagnostic protein #26

(R,K){20,20}

(K){20}

24: QMKSI KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

25: MKSIK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

26: KSIKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

27: SIKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

28: IKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

29: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKF

(K){20}

30: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKFH

(K){20}

31: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKHF

(K){20}

32: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKHFL

(K){20}

33: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKFHL

ABG26721 ck: 287 len: 95 i Abg26721 Novel human diagnostic protein #26

(R,K){20,20}

(K){20}

31: EKEKE KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

32: KEKEK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

33: EKEKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

34: KEKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

35: EKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

36: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

37: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

38: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

39: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

40: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

41: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

1

ABG26722 ck: 4831 len: 127 i Abg26722 Novel human diagnostic protein #26

(R,K){20,20}

(K){20}

44: KKKKE KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

45: KKEEK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

46: KEKKK KKKKKKKKKKKKKKKKKKKKKKKKKKE

(K){20}

47: EKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKE

(K){20}

48: EKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKE

(K){20}

49: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKE

(K){20}

87: EKEEE KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

88: KEEEK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

89: EEEKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

90: EKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K){20}

```

91: EKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
92: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
89: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
93: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
94: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
95: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
96: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
97: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
98: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
99: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
100: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
101: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
102: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
103: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
104: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
105: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
106: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KK
      (K) {20}
107: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK K
      (K) {20}
108: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK
      (K) {20}

ABg26723 ck: 7054 len: 98 ! Abg26723 Novel human diagnostic protein #26
      (R,K){20..20}
42: EKEKE KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
43: KEKEK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
44: EKEKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
45: KEKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
46: EKKKK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}

```

[illegible]

1

(R,K){20,20}
(K){20}
259: KEEEE KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
260: EEEEEK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
261: EEEEEK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
262: EEEEEK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
263: EKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
264: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
265: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
266: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
267: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
268: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
269: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
270: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
271: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
272: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
273: KKKKK KKKKKKKKKKKKKKKKKKK KKK
(K){20}
274: KKKKK KKKKKKKKKKKKKKKKKKK KK
(K){20}
275: KKKKK KKKKKKKKKKKKKKKKKKK K
(K){20}
276: KKKKK KKKKKKKKKKKKKKKKKKK

ABG26725 ck: 909 len: 131 ! Abg26725 Novel human diagnostic protein #26

(R,K){20,20}
(R,K){20}
70: KKKKE KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
71: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
72: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
73: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

1

74: EKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
75: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
76: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
77: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
78: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
79: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
80: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
81: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
82: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
83: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

ABG26726 ck: 9300 len: 677 ! Abg26726 Novel human diagnostic protein #26

(R,K){20,20}
(R,K){20}
240: EKEKE RRRRRRRRRRRRRRRRRR KKKKK
(R,K){20}
241: KEKER KRRRRRRRRRRRRRRRRR KKKKK
(R,K){20}
242: EKERK RRRRRRRRRRRRRRRRRR KKKKK
(R,K){20}
243: KERKR KRRRRRRRRRRRRRRRRR KKKKK
(R,K){20}
244: ERRKR RRRRRRRRRRRRRRRRRR KKKKK
(R,K){20}
245: RKRKR KRRRRRRRRRRRRRRRRR KKKKK
(R,K){20}
246: KRRKR RRRRRRRRRRRRRRRRRR KKKKK
(R,K){20}
247: RKRKR KRRRRRRRRRRRRRRRRR KKKKK
(R,K){20}
248: KRRKR RRRRRRRRRRRRRRRRRR KKKKK
(R,K){20}
249: RKRKR KRRRRRRRRRRRRRRRRR KKKKK
(K){20}
250: KRRKR KRRRRRRRRRRRRRRRRR KKKKK
(K){20}
251: RKRKR KRRRRRRRRRRRRRRRRR KKKKK
(K){20}
252: KRRKR KRRRRRRRRRRRRRRRRR KKKKK
(K){20}

253: RKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 254: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 255: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 256: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 257: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 258: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 259: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 260: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R, K) {20}
 261: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R, K) {20}
 262: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R, K) {20}
 263: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R, K) {20}
 264: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R, K) {20}
 265: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R, K) {20}

ABG26727 ck: 813 len: 329 ! Abg26727 Novel human diagnostic protein #26

215: EKEKE KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R, K) {20, 20}
 (K) {20}
 216: KEKEK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 217: EKEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 218: KEKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 219: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 220: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 221: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 222: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 223: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 224: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}

225: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 226: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 227: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 228: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 229: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 230: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}

ABG26879 ck: 2669 len: 800 ! Abg26879 Novel human diagnostic protein #26

559: EREGE KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R, K) {20, 20}
 (R, K) {20}
 560: REQEK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R, K) {20}
 561: EQEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R, K) {20}
 562: QEKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 563: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 564: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}

ABG28885 ck: 8668 len: 137 ! Abg28885 Novel human diagnostic protein #28

35: EEEEG RRRRRRRRRRRRRRRRRRRR RRRKK
 (R, K) {20, 20}
 (R) {20}
 36: EEEGR RRRRRRRRRRRRRRRRRRRR RRRKK
 (R) {20}
 37: EEEGR RRRRRRRRRRRRRRRRRRRR RRRKK
 (R) {20}
 38: EGGRR RRRRRRRRRRRRRRRRRRRR RRRKK
 (R) {20}
 39: GRRRR RRRRRRRRRRRRRRRRRRRR RRRKK
 (R) {20}
 40: RRRRR RRRRRRRRRRRRRRRRRRRR RRRKK
 (R, K) {20}
 41: RRRRR RRRRRRRRRRRRRRRRRRRR RRRKK
 (R, K) {20}

AAG65985 ck: 8085 len: 154 ! Aag65985 B726P splice variant sequence. 2/2

114: TQLRQ KKKKKKKKKKKKKKKKKKKKK KKKKK
 (R, K) {20, 20}
 (K) {20}
 115: QLROK KKKKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}

```

16: LROKK KKKKKKKKKKKKKKKKKKKK KKKK      (K) {20}
17: ROKKK KKKKKKKKKKKKKKKKKKKK KKKK      (K) {20}
18: OKKKK KKKKKKKKKKKKKKKKKKKK KKKK      (K) {20}
19: KKKKK KKKKKKKKKKKKKKKKKKKK KKKK      (K) {20}
20: KKKKK KKKKKKKKKKKKKKKKKKKK KKKK      (K) {20}
21: KKKKK KKKKKKKKKKKKKKKKKKKK KKKK      (K) {20}
22: KKKKK KKKKKKKKKKKKKKKKKKKK KKKK      (K) {20}
23: KKKKK KKKKKKKKKKKKKKKKKKKK KKKK      (K) {20}
24: RKKKK KKKKKKKKKKKRRRKRKR RRKR      (R,K) {20}

ABB27693 ck: 5383 len: 86 ! Abb27693 Human peptide #544 encoded by break
(R,K) {20,20}
(R,X) {20}
15: RRRRG RRRRRKKKKKKKKKKKKK KRKR      (R,K) {20}
16: RRRGR RRRKKKKKKKKKKKKK KRKR      (R,K) {20}
17: RRRGR RRRKKKKKKKKKKKKK RRRR      (R,K) {20}
18: RGRRR RRRKKKKKKKKKKKKK RRRR      (R,K) {20}
19: GRRRR RRRKKKKKKKKKKKKK RRRR      (R,K) {20}
20: RRRRR KKKKKKKKKKKKKKKK RRRR      (R,K) {20}
21: RRRRK KKKKKKKKKKKKKKKK RRRR      (R,K) {20}
22: RRRKK KKKKKKKKKKKKKKKK RRRR      (R,K) {20}
23: RRRKK KKKKKKKKKKKKKKKK RRRR      (R,K) {20}
24: RKKKK KKRRKKKKKKRRRKRKR RRKR      (R,K) {20}

```

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25: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
26: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
27: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
28: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
29: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
30: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
31: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
32: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
33: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
34: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
35: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
36: KKKK (R,K){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
37: KKKK (R){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
38: KKKK (R){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
39: KKKK (R){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
40: KKKK (R){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
41: KKKK (R){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
42: KKKK (R){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
43: KKKK (R){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
44: KKKK (R){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR
45: KKKK (R){20} KKKKKKKKKKKKKKKKKKKRRRRR RRRR

AB28750 ck: 1334 len: 86 ! Ab28750 Peptide #1401 encoded by breast ce
(R,K){20,20}
(R,K){20}
EEEEG RRRKKKKKKKKKKKKKKKKKK KKKK
(R,K){20}
58: EEEGR RRRKKKKKKKKKKKKKKKKKK KKKK

```

59: EEGRR $(R,K)\{20\}$ RKKKKKKKKKKKKKKKKKK
60: EGRRR $(R,K)\{20\}$ KKKKKKKKKKKKKKKKKKK
61: GRRRR $(R,K)\{20\}$ KKKKKKKKKKKKKKKKKKK
62: RRRKK $(R,K)\{20\}$ KKKKKKKKKKKKKKKKKKK
63: RRRKK $(R,K)\{20\}$ KKKKKKKKKKKKKKKKKKK
64: RKKKK $(R,K)\{20\}$ KKKKKKKKKKKKKKKKKKK
65: KKKRR $(R,K)\{20\}$ KKKKKKKKKKKKKKKKKKK
66: KKKRK $(R,K)\{20\}$ KKKKKKKKKKKKKKKKK K
67: KKKRR $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
1
ABB28840 ck: 9082 len: 167 ! Abb28840 Peptide #1491 encoded by breast ce
33: EEEGR $(R,K)\{20,20\}$ RRRRRRRRRRRRRRRRRRR
34: EGRGR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
35: GRRRR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
1
ABB29360 ck: 2276 len: 89 ! Abb29360 Peptide #2011 encoded by breast ce
23: EEEEE $(R,K)\{20,20\}$ KKKKKKKKKKKKKKKKKKK
46: KEEEE $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
47: KEEEE $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
48: EEEKK $(R,K)\{20\}$ KKKKKKKKKKKKKKKKKKK
49: EEEKK $(R,K)\{20\}$ KKKKKKKKKKKKKKKKKKK
1
ABB29645 ck: 1939 len: 130 ! Abb29645 Peptide #2296 encoded by breast ce
42: EGRKE $(R,K)\{20,20\}$ RRRRRRRRRRRRRRRRRRR
43: GRKER $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
44: RKERR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR

45: KERRR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
46: EERRR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
47: RRRRR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
48: RRRRR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
49: RRRRR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
50: RRRRR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
51: RRRRR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
52: RRRRR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
53: RRRRR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
54: RRRRR $(R,K)\{20\}$ RRRRRRRRRRRRRRRRRRR
1
ABB29928 ck: 3607 len: 88 ! Abb29928 Peptide #2579 encoded by breast ce
39: EKKRE $(R,K)\{20,20\}$ KKKKKKKKKKKKKKKKKKK
40: RKREK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
41: KREKK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
42: REKKK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
43: EKKKK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
44: KKKKK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
45: KKKKK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
46: KKKKK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
47: KKKKK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
48: KKKKK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
49: KKKKK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
50: KKKKK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK
51: KKKKK $(K)\{20\}$ KKKKKKKKKKKKKKKKKKK

52: KKKK (K) {20} KKKK
53: KKKK (K) {20} KKKK
54: KKKK (K) {20} KKKK
55: KKKK (K) {20} KKKK
56: KKKK (K) {20} KKKK
57: KKKK (K) {20} KKKK
58: KKKK (K) {20} KKKK
59: KKKK (K) {20} KKKK
60: KKKK (K) {20} KKKK
61: KKKK (K) {20} KKKK
62: KKKK (K) {20} KKKK
63: KKKK (K) {20} KKKK
64: KKKK (K) {20} KKKK
65: KKKK (K) {20} KKKK
66: KKKK (K) {20} KKKK
67: KKKK (K) {20} KKKK
68: KKKK (K) {20} K
69: KKKK (K) {20} KKKK

1
ABB30512 ck: 3937 len: 85 ! Abb30512 Peptide #3163 encoded by breast ce
(R,K) {20,20}
(K) {20}

1: KKKK
2: K (K) {20} KKKK
3: KK (K) {20} KKKK
4: KKK (K) {20} KKKK
5: KKKK (K) {20} KKKK

6: KKKK (K) {20} KKKK
7: KKKK (K) {20} KKKK
8: KKKK (K) {20} KKKK
9: KKKK (K) {20} KKKK
10: KKKK (K) {20} KKKK
11: KKKK (K) {20} KKKK

1
ABB32308 ck: 1560 len: 88 ! Abb32308 Peptide #4959 encoded by breast ce
(R,K) {20,20}
(R) {20}

43: RRRG (R) {20} RRRR
44: RRRG (R) {20} RRRR
45: RRRG (R) {20} RRRR
46: RRRG (R) {20} RRRR
47: RRRG (R) {20} RRRR
48: RRRG (R) {20} RRRR
49: RRRG (R) {20} RRRR
50: RRRG (R) {20} RRRR
51: RRRG (R) {20} RRRR
52: RRRG (R) {20} RRRR
53: RRRG (R) {20} RRRR
54: RRRG (R) {20} RRRR
55: RRRG (R) {20} RRRR
56: RRRG (R) {20} RRRR
57: RRRG (R) {20} RRRR
58: RRRG (R) {20} RRRR
59: RRRG (R) {20} RRRR

1

60: RRRR (R, K) {20} RRRRRRRRRRRRRRRRRR RRVTN
61: RRRR (R, K) {20} RRRRRRRRRRRRRRRRRR RVNTN
62: RRRR (R, K) {20} RRRRRRRRRRRRRRRRRR NTVNE
ABB33064 ck: 5383 len: 86 1 Abb33064 Peptide #570 encoded by human foet
(R, K) {20, 20}
15: RRRG RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
16: RRRG RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
17: RRRG RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
18: RRRG RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
19: RRRG RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
20: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
21: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
22: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
23: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
24: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
25: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
26: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
27: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
28: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
29: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
30: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
31: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
32: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
33: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
34: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}

1

35: RRRR (R, K) {20} RRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}
36: RRRR RRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}
37: RRRR RRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}
38: RRRR RRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}
39: RRRR RRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}
40: RRRR RRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}
41: RRRR RRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}
42: RRRR RRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}
43: RRRR RRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}
44: RRRR RRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}
45: RRRR RRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}
ABB33937 ck: 1334 len: 86 1 Abb33937 Peptide #1443 encoded by human foet
(R, K) {20, 20}
57: EEEG RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
58: EEEG RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
59: EEEG RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
60: EEEG RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
61: GRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
62: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
63: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
64: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
65: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
66: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
67: RRRR RRRRRRRRRRRRRRRRRR KRRR
(R, K) {20}
ABB34024 ck: 9082 len: 167 1 Abb34024 Peptide #1530 encoded by human foet

1

(R,K){20,20}
33: EEGRG RRRRRRRRRRRRRRRRRR RRGGR
(R,K){20}
34: EGRGR RRRRRRRRRRRRRRRRRR RGGGR
(R,K){20}
35: GGRGR RRRRRRRRRRRRRRRRRR GGGR

1

ABB34533 ck: 2276 len: 89 ! Abb34533 Peptide #2039 encoded by human foe
(R,K){20,20}
23: EEEEE KKKKKKKKKKKKKKKKKKK EEEK
(R,K){20}
46: KKEEE KKKKKKKKKKKKKKKKKKK KRKEE
(K){20}
47: KEEK KKKKKKKKKKKKKKKKKKK RKEE
(R,K){20}
48: EEEK KKKKKKKKKKKKKKKKKKK KESEE
(R,K){20}
49: EEEK KKKKKKKKKKKKKKKKKKK EEEEE

1

ABB34819 ck: 1939 len: 130 ! Abb34819 Peptide #2325 encoded by human foe
(R,K){20,20}
42: EGRKE RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
43: GRKER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
44: RKEER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
45: KERER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
46: ERRER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
47: RRRER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
48: RRRER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
49: RRRER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
50: RRRER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
51: RRRER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
52: RRRER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
53: RRRER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}
54: RRRER RRRRRRRRRRRRRRRRRR RRRR
(R,K){20}

1

ABB35110 ck: 3607 len: 88 ! Abb35110 Peptide #2616 encoded by human foe
(R,K){20,20}
39: ERKRE KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
40: RKREK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
41: KRERK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
42: REKRE KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
43: ERKRE KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
44: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
45: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
46: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
47: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
48: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
49: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
50: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
51: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
52: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
53: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
54: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
55: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
56: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
57: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
58: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
59: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
60: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
61: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}

62: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
63: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
64: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
65: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
66: KKKKK KKKKKKKKKKKKKKKKKKK KKK
(K) {20}
67: KKKKK KKKKKKKKKKKKKKKKKKK KK
(K) {20}
68: KKKKK KKKKKKKKKKKKKKKKKKK K
(K) {20}
69: KKKKK KKKKKKKKKKKKKKKKKKK

ABB35676 ck: 3937 len: 85 ! Abb35676 Peptide #3182 encoded by human foe

1

(R, K) {20, 20}
(K) {20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
2: K KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: KK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKK EEEEE

ABB36406 ck: 2686 len: 71 ! Abb36406 Peptide #3912 encoded by human foe

1

(R, K) {20, 20}
(K) {20}
20: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
21: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
22: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

(K) {20}
23: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
24: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
25: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
26: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
27: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
28: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
29: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
30: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
31: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
32: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
33: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
34: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
35: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
36: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
37: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
38: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
39: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
40: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
41: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
42: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
43: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
44: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
45: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R, K) {20}
46: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

47: KKKKK KKKKKKKKKKKKKKKKK KKSAN
(K) {20}
48: KKKKK KKKKKKKKKKKKKKKKK KSAN
(K) {20}
49: KKKKK KKKKKKKKKKKKKKKKK SAN

1

ABB37567 ck: 1560 len: 88 1 Abb37567 Peptide #5073 encoded by human foe

(R,K) {20,20}

(R) {20}

43: RRRRG RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

44: ERRGR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

45: RRGRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

46: RGRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

47: GRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

48: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

49: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

50: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

51: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

52: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

53: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

54: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

55: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

56: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

57: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

58: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

59: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

60: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

61: RRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

62: RRRRR RRRRRRRRRRRRRRRRR RRRRR

1

ABB37780 ck: 2324 len: 36 1 Abb37780 Peptide #5286 encoded by human foe

(R,K) {20,20}

(R,K) {20}

16: KERKT KKKRRRRRRRRRRRRRR R

(R,K) {20}

17: ERKTX KKKRRRRRRRRRRRRRR

1

ABB40272 ck: 8343 len: 66 1 Abb40272 Peptide #7778 encoded by human foe

(R,K) {20,20}

(R,K) {20}

6: ETERE KKKRRRRRRRRRRRRRR KKKKK

7: TEREK KKKRRRRRRRRRRRRRR KKKKK

(R,K) {20}

8: EREKK KKKRRRRRRRRRRRRRR KKKKK

(R,K) {20}

9: REKKK KKKRRRRRRRRRRRRRR KKKKK

1

ABB42642 ck: 2394 len: 57 1 Abb42642 Peptide #10148 encoded by human foe

(R,K) {20,20}

(R) {20}

20: EEEEG RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

21: EEEGR RRRRRRRRRRRRRRRRR RRRRG

(R) {20}

22: EEEGR RRRRRRRRRRRRRRRRR RRRGR

(R) {20}

23: EGGRR RRRRRRRRRRRRRRRRR RRRGR

(R) {20}

24: GRRRR RRRRRRRRRRRRRRRRR RRRRR

(R) {20}

25: RRRRR RRRRRRRRRRRRRRRRR GRRRR

1

ABB43181 ck: 4228 len: 24 1 Abb43181 Peptide #10687 encoded by human foe

(R,K) {20,20}

(R,K) {20}

1: RRRRRRRRRRRRRRRRRRR RRRRT

(R,K) {20}

2: R RRRRRRRRRRRRRRRRR RRT

(R,K) {20}

3: RR RRRRRRRRRRRRRRRRR RT

(R,K) {20}

4: RRR RRRRRRRRRRRRRRRRR T

1

ABB44317 ck: 4695 len: 51 1 Abb44317 Peptide #11823 encoded by human foe

(R,K) {20,20}

(R,K) {20}

18: LFKPM KKKRRRRRRRRRRRRRR KKLTT

(R,K) {20}

19: FKPMR KRRRRRRRRRRRRRRRR KLT TT

20: KPMRK RRRKKRRKKKKRRRRKK LTTT
(R, K) {20}

ABBI7165 ck: 8887 len: 42 1 Abbi7165 Human nervous system related poly

(R, K) {20, 20}

21: FTTTE KKKKKKKKKKKKKKKKKKK KX
(K) {20}

22: FTTEK KKKKKKKKKKKKKKKKKKK X
(K) {20}

ABBI8534 ck: 5383 len: 86 1 Abbi8534 Protein #533 encoded by probe for

(R, K) {20, 20}

15: RRRRG RRRRRKKKKKKKKKKKKKK KRRR
(R, K) {20}

16: RRRGR RRRKKKKKKKKKKKKKKKK KRRR
(R, K) {20}

17: RRGRR RRRKKKKKKKKKKKKKKKK RRRR
(R, K) {20}

18: RGRRR RRRKKKKKKKKKKKKKKKK RRRR
(R, K) {20}

19: GRRRR RKKKKKKKKKKKKKKKKKK RRRR
(R, K) {20}

20: RRRRR KKKKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

21: RRRRK KKKKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

22: RRRKK KKKKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

23: RRRKK KKKKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

24: RKKKK KRRKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

25: KKKKK KRRKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

26: KKKKK RKKKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

27: KKKKK KKKKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

28: KKKKK KKKKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

29: KKKKK KKKKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

30: KKKKK KRRKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

31: RKKKK KRRKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

32: KKKKK RKKKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

33: KKKKK KKKKKKKKKKKKKKKKKRR RRRR
(R, K) {20}

34: KKKKK KKKRRRRRRRRRRRRRRRR RRRR
(R, K) {20}

35: KKKKK KRRRRRRRRRRRRRRRRRR RRRR
(R, K) {20}

36: KKKKK KRRRRRRRRRRRRRRRRRR RRRR
(R) {20}

37: RKKKK RRRRRRRRRRRRRRRRRRR RRRR
(R) {20}

38: KKKKK RRRRRRRRRRRRRRRRRRR RRRR
(R) {20}

39: KKKKK RRRRRRRRRRRRRRRRRRR RRRR
(R) {20}

40: KKKKK RRRRRRRRRRRRRRRRRRR RRRR
(R) {20}

41: KRRRR RRRRRRRRRRRRRRRRRRR RRRR
(R) {20}

42: RRRRR RRRRRRRRRRRRRRRRRRR RRRR
(R) {20}

43: RRRRR RRRRRRRRRRRRRRRRRRR RRRR
(R) {20}

44: RRRRR RRRRRRRRRRRRRRRRRRR RRRR
(R) {20}

45: RRRRR RRRRRRRRRRRRRRRRRRR RRRR
(R) {20}

ABBI9373 ck: 1334 len: 86 1 Abbi9373 Protein #1372 encoded by probe for

(R, K) {20, 20}

57: EEEEG RRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}

58: EEEGR RRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}

59: EEEGR RRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}

60: EEEGR RRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}

61: GRRRK KRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}

62: RRRKK KRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}

63: RRRKK KRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}

64: RRRKK KRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}

65: KKKKK KRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}

66: KKKKK RRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}

67: KRRRR KKKKKKKKKKKKKKKKKKK KKKK
(R, K) {20}

1

ABB19467 ck: 9082 len: 167 i Abb19467 Protein #1466 encoded by probe for
 (R,K){20,20}
 33: EEEGRG RRRRRRRRRRRRRRRRRRRR RRGCG
 (R,K){20}
 34: EGRGR RRRRRRRRRRRRRRRRRRRR RGGGR
 (R,K){20}
 35: GGRGR RRRRRRRRRRRRRRRRRRRR GGGRR

1

ABB19943 ck: 2276 len: 89 i Abb19943 Protein #1942 encoded by probe for
 (R,K){20,20}
 23: EEEEE KKKKKKKKKKKKKKKKKKK EEEKK
 (R,K){20}
 46: KKEEE KKKKKKKKKKKKKKKKKKK KRKEE
 (K){20}
 47: KEEKK KKKKKKKKKKKKKKKKKKK RKEE
 (R,K){20}
 48: EEEKK KKKKKKKKKKKKKKKKKKK KEEEE
 (R,K){20}
 49: EEEKK KKKKKKKKKKKKKKKKKKK EEEEE

1

ABB20238 ck: 1939 len: 130 i Abb20238 Protein #2237 encoded by probe for
 (R,K){20,20}
 42: EGRKE RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}
 43: GRKER RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}
 44: RKEER RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}
 45: KEERR RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}
 46: ERRER RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}
 47: RRRER RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}
 48: RRRER RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}
 49: RRRER RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}
 50: RRRER RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}
 51: RRRER RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}
 52: RRRER RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}
 53: RRRER RRRRRRRRRRRRRRRRRRRR RRRRR
 (R,K){20}

1

54: RRRRR RRRRRRRRRRRRRRRRRRRR EEEEE
 (R,K){20}
 ABB20531 ck: 3607 len: 88 i Abb20531 Protein #2530 encoded by probe for
 (R,K){20,20}
 39: ERKRE KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 40: RKREK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 41: KREKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 42: REKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 43: EKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 44: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 45: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 46: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 47: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 48: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 49: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 50: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 51: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 52: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 53: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 54: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 55: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 56: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 57: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 58: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 59: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K){20}
 60: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

61: KKKK (K) {20}
 62: KKKK (K) {20}
 63: KKKK (K) {20}
 64: KKKK (K) {20}
 65: KKKK (K) {20}
 66: KKKK (K) {20}
 67: KKKK (K) {20}
 68: KKKK (K) {20}
 69: KKKK (K) {20}

ABb21105 ck: 3937 len: 95 i Abb21105 Protein #3104 encoded by probe for

1

1: KKKK (R, K) {20,20}
 2: K KKKK (K) {20}
 3: KK KKKK (K) {20}
 4: KK KKKK (K) {20}
 5: KKKK (K) {20}
 6: KKKK (K) {20}
 7: KKKK (K) {20}
 8: KKKK (K) {20}
 9: KKKK (K) {20}
 10: KKKK (K) {20}
 11: KKKK (K) {20}

ABb21763 ck: 2686 len: 71 i Abb21763 Protein #3762 encoded by probe for

1

20: KKKKQ KKKK (R, K) {20,20}
 21: KKKK (K) {20}

22: KKQK (K) {20}
 23: KQKK (K) {20}
 24: QKKK (K) {20}
 25: KKKK (R, K) {20}
 26: KKKK (R, K) {20}
 27: KKKK (R, K) {20}
 28: KKKK (R, K) {20}
 29: KKKK (R, K) {20}
 30: KKKK (R, K) {20}
 31: KKKK (R, K) {20}
 32: KKKK (R, K) {20}
 33: KKKK (R, K) {20}
 34: KKKK (R, K) {20}
 35: KKKK (R, K) {20}
 36: KKKK (R, K) {20}
 37: KKKK (R, K) {20}
 38: KKKK (R, K) {20}
 39: KKKK (R, K) {20}
 40: KKKK (R, K) {20}
 41: KKKK (R, K) {20}
 42: KKKK (R, K) {20}
 43: KKKK (R, K) {20}
 44: KKKK (R, K) {20}
 45: KKKK (R, K) {20}

46: KKKKK (R,K){20} KKKSA
 47: KKKKK (K){20} KKSAA
 48: KKKKK (K){20} KSAH
 49: KKKKK (K){20} SAH

1 ABB22862 ck: 1560 len: 88 ! ABB22862 Protein #4661 encoded by probe for
 (R,K){20,20}

43: RRRRG (R){20} RRRRR
 44: RRRGR (R){20} RRRRR
 45: RRGRR (R){20} RRRRR
 46: RGRRR (R){20} RRRRR
 47: GRRRR (R){20} RRRRR
 48: RRRRR (R){20} RRRRR
 49: RRRRR (R){20} RRRRR
 50: RRRRR (R){20} RRRRR
 51: RRRRR (R){20} RRRRR
 52: RRRRR (R){20} RRRRR
 53: RRRRR (R){20} RRRRR
 54: RRRRR (R){20} RRRRR
 55: RRRRR (R){20} RRRRR
 56: RRRRR (R){20} RRRRR
 57: RRRRR (R){20} RRRRR
 58: RRRRR (R){20} RRRRR
 59: RRRRR (R){20} RRRRR
 60: RRRRR (R){20} RRRRR
 61: RRRRR (R){20} RRRRR

62: RRRRR (R){20} RRRRR NTNNE
 1 ABB23064 ck: 2324 len: 36 ! ABB23064 Protein #5063 encoded by probe for
 (R,K){20,20}
 16: KRRRT (R,K){20} RRRRR R
 17: ERRTK (R,K){20} RRRRR RRRRR

1 ABB24685 ck: 8343 len: 66 ! ABB24685 Protein #6684 encoded by probe for
 (R,K){20,20}

6: ETERE (R,K){20} KKKKK
 7: TEREK (R,K){20} KKKKK
 8: EREKK (R,K){20} KKKKK
 9: REKKK (R,K){20} KKKKK

1 ABB25988 ck: 2394 len: 57 ! ABB25988 Protein #7987 encoded by probe for
 (R,K){20,20}

20: EEEEG (R){20} RRRRR
 21: EEEGR (R){20} RRRRR
 22: EGGRR (R){20} RRRRR
 23: EGRRR (R){20} RRRRR
 24: GRRRR (R){20} RRRRR
 25: RRRRR (R){20} RRRRR

1 ABB27176 ck: 4895 len: 51 ! ABB27176 Protein #9175 encoded by probe for
 (R,K){20,20}

18: LFKPM (R,K){20} RRRRR KLLTT
 19: FKPMR (R,K){20} RRRRR KLLTT
 20: KPMRK (R,K){20} RRRRR LTTT

1 ABB10296 ck: 3983 len: 292 ! ABB10296 Human cDNA SEQ ID NO: 604. 1/2002
 (R,K){20,20}

273: QVFAP (R,K){20} RRRRR

ABB10485 ck: 7611 len: 315 ! ABB10485 Human cDNA SEQ ID NO: 793. 1/2002

1

(R,K){20,20}
 273: QVFAP RXXXXXXXXXXXXXXXXX KGGRS
 (K){20}
 274: VFAPR KXXXXXXXXXXXXXXXXX GGRSR

1

AAU21948 ck: 444 len: 66 i Aau21948 Human cardiovascular system antige
 (R,K){20,20}
 35: SMTFS KXXXXXXXXXXXXXXXXX KXGKK
 (K){20}
 36: MTFSS KXXXXXXXXXXXXXXXXX XGKKK

1

AAU22148 ck: 4665 len: 34 i Aau22148 Human cardiovascular system antige
 (R,K){20,20}
 10: PELLK KXXXXXXXXXXXXXXXXX KKKKK
 (K){20}
 11: ELLLK KXXXXXXXXXXXXXXXXX KKKK
 (K){20}
 12: LLLKK KXXXXXXXXXXXXXXXXX KKK
 (K){20}
 13: LLKKK KXXXXXXXXXXXXXXXXX KK
 (K){20}
 14: LKKKK KXXXXXXXXXXXXXXXXX K
 (K){20}
 15: KKKKK KXXXXXXXXXXXXXXXXX

1

AAU22186 ck: 269 len: 76 i Aau22186 Human cardiovascular system antige
 (R,K){20,20}
 37: TPSRA KXXXXXXXXXXXXXXXXX KKKKK
 (K){20}
 38: PSRAK KXXXXXXXXXXXXXXXXX KKKKK
 (K){20}
 39: SRAKK KXXXXXXXXXXXXXXXXX KKKKK
 (K){20}
 40: RAKKK KXXXXXXXXXXXXXXXXX KKKKK
 (K){20}
 41: AKKKK KXXXXXXXXXXXXXXXXX KKKKK
 (K){20}
 42: KKKKK KXXXXXXXXXXXXXXXXX KKKKK
 (K){20}
 43: KKKKK KXXXXXXXXXXXXXXXXX KKKKI
 (K){20}
 44: KKKKK KXXXXXXXXXXXXXXXXX XKIK

1

AAU22374 ck: 8278 len: 53 i Aau22374 Human cardiovascular system antige
 (R,K){20,20}
 (K){20}

30: NCGIL KXXXXXXXXXXXXXXXXX KKKK

31: CGILK KXXXXXXXXXXXXXXXXX KKK

32: GILKK KXXXXXXXXXXXXXXXXX KK

33: ILKKK KXXXXXXXXXXXXXXXXX K

34: LKKKK KXXXXXXXXXXXXXXXXX

1

AAU23799 ck: 6158 len: 272 i Aau23799 Novel human enzyme polypeptide #88:
 (R,K){20,20}
 238: SPANA KXXXXXXXXXXXXXXXXX KKKKG
 (K){20}
 239: PANAK KXXXXXXXXXXXXXXXXX KKKGR
 (K){20}
 240: ANAKK KXXXXXXXXXXXXXXXXX KKGRR
 (K){20}
 241: NAKKK KXXXXXXXXXXXXXXXXX KGRPX
 (K){20}
 242: AKKKK KXXXXXXXXXXXXXXXXX GRPXX

1

AAU27944 ck: 1121 len: 69 i Aau27944 Human config polypeptide sequence
 (R,K){20,20}
 40: VPPLT RXXXXXXXXXXXXXXXXX KKKKK
 (K){20}
 41: PPLTR KXXXXXXXXXXXXXXXXX KKKKK
 (K){20}
 42: PLTRK KXXXXXXXXXXXXXXXXX KKKKR
 (K){20}
 43: LTRKK KXXXXXXXXXXXXXXXXX KKKRG
 (K){20}
 44: TRKKK KXXXXXXXXXXXXXXXXX KKRGA
 (K){20}
 45: RKKKK KXXXXXXXXXXXXXXXXX KRGA
 (K){20}
 46: KKKKK KXXXXXXXXXXXXXXXXX RGAL
 (R,K){20}
 47: KKKKK KXXXXXXXXXXXXXXXXX GAL

1

AAU31467 ck: 4264 len: 657 i Aau31467 Novel human secreted protein #1958
 (R,K){20,20}
 19: RRRRP RRRRRRRRRRRRRRRRRR RRRRL
 (R){20}
 20: RRRPR RRRRRRRRRRRRRRRRRR RRRLG
 (R){20}
 21: RRRPR RRRRRRRRRRRRRRRRRR RRLGL

22: RPRRR RRRRRRRRRRRRRRRR RLGLE
(R) {20}
23: PRRRR RRRRRRRRRRRRRRRR LGLE
(R) {20}

1 AaU3348 ck: 8085 len: 154 ! AaU3348 Human breast cancer protein encode

(R,K) {20,20}
(K) {20}

114: TQLRQ KKKKKKKKKKKKKKKKK KKKK

(K) {20}

115: QLRQK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

116: LRQK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

117: RQK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

118: QKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

119: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

120: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

121: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

122: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

123: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

124: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

125: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

126: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

127: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

128: KKKK KKKKKKKKKKKKKKKKK KKKK

AaE09664 ck: 1663 len: 87 ! AaE09664 Human pancreatic related protein H

(R,K) {20,20}
(K) {20}

36: KWSXK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

37: WSSXK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

38: SSXK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

39: SXK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

40: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

41: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

42: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

43: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

44: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

45: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

46: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

47: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

48: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

49: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

50: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

51: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

52: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

53: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

54: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

55: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

56: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

57: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

58: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

59: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

60: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

61: KKKK KKKKKKKKKKKKKKKKK KKKK

(K) {20}

62: KKKK KKKKKKKKKKKKKKKKK KKKK

(R,K) {20}

63: KKKK KKKKKKKKKKKKKKKKK KKKK

AaM95365 ck: 5626 len: 139 ! AaM95365 Human reproductive system related

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1      (R,K){20,20}
111: IHLNL KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
112: HLNLK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
113: LNLKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
114: NLKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

AAM96607 ck: 4751 len: 80   ! Aam96607 Human reproductive system related
      (R,K){20,20}
      (K){20}
61: KKKFD KKKKKKKKKKKKKKKKKKKKK

AAU18162 ck: 7907 len: 39   ! Aau18162 Novel human DNA-binding protein #9
      (R,K){20,20}
      (K){20}
9: YFEDL KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
10: FEDLK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
11: EDLKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
12: DLKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
13: LKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
14: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

AAU18167 ck: 9194 len: 87   ! Aau18167 Novel human DNA-binding protein #1
      (R,K){20,20}
      (K){20}
52: KIILL KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
53: IILLK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
54: ILLKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
55: LLKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
56: LKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
57: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

AAU18168 ck: 8659 len: 104   ! Aau18168 Novel human DNA-binding protein #1
      (R,K){20,20}
      (K){20}
75: PLGGQ KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}

```

```

76: LGGQK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
77: GGQKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
78: GQKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
79: QKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
80: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

AAU18171 ck: 9398 len: 48   ! Aau18171 Novel human DNA-binding protein #1f
      (R,K){20,20}
      (K){20}
2: Q KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
3: QK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
4: QKK KKKKKKKKKKKKKKKKKKKKK KKKKK

AAU18177 ck: 8278 len: 53   ! Aau18177 Novel human DNA-binding protein #2f
      (R,K){20,20}
      (K){20}
30: NCGIL KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
31: CGILK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
32: GILKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
33: ILKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
34: LKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

AAU18178 ck: 444 len: 66   ! Aau18178 Novel human DNA-binding protein #2
      (R,K){20,20}
      (K){20}
35: SMTFS KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
36: MTFSS KKKKKKKKKKKKKKKKKKKKK KKKKK

AAU18179 ck: 5503 len: 50   ! Aau18179 Novel human DNA-binding protein #2
      (R,K){20,20}
      (K){20}
30: IICLL KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
31: ICLLK KKKKKKKKKKKKKKKKKKKKK KKKKK

AAU18184 ck: 5691 len: 108   ! Aau18184 Novel human DNA-binding protein #3
      (R,K){20,20}
      (K){20}
78: VRPCL KKKKKKKKKKKKKKKKKKKKK KKKKK

```


79: RPCLK KKKKKKKKKKKKKKKKKKK (K) {20}
 80: PCLKK KKKKKKKKKKKKKKKKKKK (K) {20}
 81: CLKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 82: LKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 83: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 84: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 85: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 86: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 87: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 88: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 89: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}

AAU18192 ck: 6029 len: 63 ! Aau18192 Novel human DNA-binding protein #3
 (R, K) {20, 20}
 (K) {20}

40: KLTLK KKKKKKKKKKKKKKKKKKK ISWG

AAU18200 ck: 7170 len: 63 ! Aau18200 Novel human DNA-binding protein #4
 (R, K) {20, 20}
 (K) {20}

37: TPSRA KKKKKKKKKKKKKKKKKKK (K) {20}
 38: PSRAK KKKKKKKKKKKKKKKKKKK (K) {20}
 39: SRAKK KKKKKKKKKKKKKKKKKKK (K) {20}
 40: RAKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 41: AKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 42: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 43: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 44: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}

AAU18204 ck: 6110 len: 61 ! Aau18204 Novel human DNA-binding protein #5
 (R, K) {20, 20}
 (K) {20}

28: RPTRP KKKKKKKKKKKKKKKKKKK (K) {20}
 29: PTPRP KKKKKKKKKKKKKKKKKKK (K) {20}
 30: TRPKK KKKKKKKKKKKKKKKKKKK (K) {20}
 31: RPKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 32: PKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 33: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 34: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 35: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 36: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 37: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}

AAU18205 ck: 5764 len: 74 ! Aau18205 Novel human DNA-binding protein #5
 (R, K) {20, 20}
 (K) {20}

40: EFLSA KKKKKKKKKKKKKKKKKKK (K) {20}
 41: FLSAK KKKKKKKKKKKKKKKKKKK (K) {20}
 42: LSARK KKKKKKKKKKKKKKKKKKK (K) {20}
 43: SAKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 44: AKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 45: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 46: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 47: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 48: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 49: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 50: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 51: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
 52: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}

53: KKKKK KKKKKKKKKKKKKKKKKKKKK KK

(K) {20}
54: KKKKK KKKKKKKKKKKKKKKKKKKKK X

AAU18206 ck: 9217 len: 68 1 Aau18206 Novel human DNA-binding protein #5

(R,K) {20,20}

38: FLEPE KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
39: LPPEK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
40: FPEKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
41: PEKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
42: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
43: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
44: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
45: KKKKK KKKKKKKKKKKKKKKKKKKKK GKXX

AAU18208 ck: 8162 len: 79 1 Aau18208 Novel human DNA-binding protein #5

(R,K) {20,20}

41: VRPRV RKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
42: RPRVR KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
43: PRVRK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
44: RVRKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
45: VRKKK KKKKKKKKKKKKKKKKKKKKK GGRFR

AAU18237 ck: 285 len: 118 1 Aau18237 Novel human DNA-binding protein #8

(R,K) {20,20}

98: EKHKQ KKKKKKKKKKKKKKKKKKKKKR G

AAU18238 ck: 5509 len: 58 1 Aau18238 Novel human DNA-binding protein #8

(R,K) {20,20}

36: FYFVC KKKKKKKKKKKKKKKKKKKKK KKK

(K) {20}
37: YFVCK KKKKKKKKKKKKKKKKKKKKK KK

(K) {20}
38: FVCKK KKKKKKKKKKKKKKKKKKKKK K

(K) {20}

39: VCKKK KKKKKKKKKKKKKKKKKKKKK

AAU18239 ck: 9074 len: 66 1 Aau18239 Novel human DNA-binding protein #8

(R,K) {20,20}

40: LVQCE KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
41: VQCEK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
42: QCEKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
43: CEKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
44: EKKKK KKKKKKKKKKKKKKKKKKKKK KKK

(K) {20}
45: KKKKK KKKKKKKKKKKKKKKKKKKKK KK

(K) {20}
46: KKKKK KKKKKKKKKKKKKKKKKKKKK K

(K) {20}
47: KKKKK KKKKKKKKKKKKKKKKKKKKK

AAU18240 ck: 8528 len: 150 1 Aau18240 Novel human DNA-binding protein #8

(R,K) {20,20}

113: SRNTV KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
114: RNTVK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
115: NTVRK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
116: TVRKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
117: VRKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
118: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
119: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
120: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
121: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
122: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
123: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

AAU18241 ck: 7676 len: 156 1 Aau18241 Novel human DNA-binding protein #8

(R,K) {20,20}

(K) {20}
108: KTTWI KKKKKKKKKKKKKKKKKKKKK KKKKK

109: TWIWK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
110: TWIWK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
111: WIKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
112: IKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
113: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
114: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
115: KKKKK KKKKKKKKKKKKKKKKKKK (R,K) {20} KKKKK

AAU18242 ck: 1736 len: 40 i Aau18242 Novel human DNA-binding protein #8

(R,K) {20,20}

(K) {20}

18: LPSLK KKKKKKKKKKKKKKKKKKK (K) {20} KKK

19: PGS�K KKKKKKKKKKKKKKKKKKK (K) {20} KKK

20: GS�KK KKKKKKKKKKKKKKKKKKK (K) {20} K

21: SLKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

AAU18244 ck: 1109 len: 98 i Aau18244 Novel human DNA-binding protein #9

(R,K) {20,20}

(K) {20}

53: OTNNT KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

54: TKNTK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

55: KNTKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

56: NTYKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

57: TKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

58: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

59: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

60: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

61: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

62: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
(K) {20}

63: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
64: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
65: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
66: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
67: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK
68: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

AAU18246 ck: 8102 len: 111 i Aau18246 Novel human DNA-binding protein #9

(R,K) {20,20}

(K) {20}

78: EFHIL KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

79: PHILK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

80: HILKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

81: ILKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

82: LKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

83: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

84: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

85: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

86: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

87: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

88: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

89: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

90: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

91: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} K

92: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

AAU18247 ck: 8102 len: 111 i Aau18247 Novel human DNA-binding protein #9
(R,K) {20,20}
(K) {20}

78: EFHIL KKKKKKKKKKKKKKKKKKK KKKKK

79: PHILK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
80: HILKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
81: ILKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
82: LKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
83: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
84: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
85: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
86: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
87: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
88: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
89: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
90: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
91: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
92: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}

AAU18248 ck: 8319 len: 53 ! Aau18248 Novel human DNA-binding protein #9
(R,K) {20,20}
(K) {20}

13: RYFKP KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
14: YFKPK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
15: FKPKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
16: KPKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
17: PKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
18: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
19: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
20: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
21: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}

22: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
23: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
24: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
25: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
26: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
27: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
28: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
29: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
30: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}

AAU18250 ck: 7918 len: 80 ! Aau18250 Novel human DNA-binding protein #9
(R,K) {20,20}
(K) {20}

50: NVLTV KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
51: VLTVK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
52: LTVKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
53: TVKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
54: VKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
55: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
56: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}

AAU18252 ck: 4882 len: 41 ! Aau18252 Novel human DNA-binding protein #9
(R,K) {20,20}
(K) {20}

8: FYCFP KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
9: YCFPK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
10: CFPKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
11: PPKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
12: PKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK (K) {20}

14: KKKK (K) {20}
15: KKKK (K) {20}
16: KKKK (K) {20}
17: KKKK (K) {20}
18: KKKK (K) {20}
19: KKKK (K) {20}
20: KKKK (K) {20}

AAU18253 ck: 5469 len: 63 ! Aau18253 Novel human DNA-binding protein #1

(R,K) {20,20}

(K) {20}

30: ITCL KKKK (K) {20}

31: ICLL KKKK (K) {20}

32: CLLK KKKK (K) {20}

33: LCLK KKKK (K) {20}

34: LKKK KKKK (K) {20}

35: KKKK KKKK (K) {20}

36: KKKK KKKK (K) {20}

37: KKKK KKKK (K) {20}

38: KKKK KKKK (K) {20}

39: KKKK KKKK (K) {20}

40: KKKK KKKK (K) {20}

41: KKKK KKKK (K) {20}

42: KKKK KKKK (K) {20}

AAU18254 ck: 5075 len: 52 ! Aau18254 Novel human DNA-binding protein #1

(R,K) {20,20}

(K) {20}

30: FIVK KKKK (K) {20}

(K) {20}

31: IVVK KKKK (K) {20}
32: VVKK KKKK (K) {20}
33: VKKK KKKK (K) {20}

AAU18255 ck: 5741 len: 47 ! Aau18255 Novel human DNA-binding protein #1

(R,K) {20,20}

(K) {20}

20: ILTF KKKK (K) {20}

21: LTFK KKKK (K) {20}

22: TTFK KKKK (K) {20}

23: TFKK KKKK (K) {20}

24: FKKK KKKK (K) {20}

25: KKKK KKKK (K) {20}

26: KKKK KKKK (K) {20}

27: KKKK KKKK (K) {20}

AAU18256 ck: 2868 len: 84 ! Aau18256 Novel human DNA-binding protein #1

(R,K) {20,20}

(K) {20}

53: KCTE KKKK (K) {20}

54: CTYE KKKK (K) {20}

55: TYEK KKKK (K) {20}

56: YEKK KKKK (K) {20}

57: EKKK KKKK (K) {20}

58: KKKK KKKK (K) {20}

59: KKKK KKKK (K) {20}

60: KKKK KKKK (K) {20}

61: KKKK KKKK (K) {20}

62: KKKK KKKK (K) {20}

AAU18257 ck: 4686 len: 73 ! Aau18257 Novel human DNA-binding protein #1

1
 (R,K){20,20}
 (K){20}
41: YLKE KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
42: LKKE KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
43: KKEK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
44: KEKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
45: EKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
46: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
47: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
48: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
49: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
50: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
51: KKKK KKKKKKKKKKKKKKKKKKK KKK
 (K){20}
52: KKKK KKKKKKKKKKKKKKKKKKK KK
 (K){20}
53: KKKK KKKKKKKKKKKKKKKKKKK K
 (K){20}
54: KKKK KKKKKKKKKKKKKKKKKKK
 (K){20}
1
AAU18258 ck: 6676 len: 74 ! Aau18258 Novel human DNA-binding protein #1
 (R,K){20,20}
 (K){20}
47: LRTQ KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
48: RTQK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
49: TQK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
50: FQK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
51: QKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
52: KKK KKKKKKKKKKKKKKKKKKK KKK
 (K){20}
1
AAU18259 ck: 2283 len: 54 ! Aau18259 Novel human DNA-binding protein #1
 (R,K){20,20}
 (K){20}
32: IVCF KKKKKKKKKKKKKKKKKKK KKK
 (K){20}

1
33: VCFK KKKKKKKKKKKKKKKKKKK KK
 (K){20}
34: PCFK KKKKKKKKKKKKKKKKKKK X
 (K){20}
1
AAU18260 ck: 7503 len: 74 ! Aau18260 Novel human DNA-binding protein #1
 (R,K){20,20}
 (K){20}
45: SHLD KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
46: HLDK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
47: LTDK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
48: TDKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
49: DKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
50: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
51: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
52: KKKK KKKKKKKKKKKKKKKKKKK KKK
 (K){20}
53: KKKK KKKKKKKKKKKKKKKKKKK KK
 (K){20}
54: KKKK KKKKKKKKKKKKKKKKKKK K
 (K){20}
55: KKKK KKKKKKKKKKKKKKKKKKK
 (K){20}
1
AAU18262 ck: 5199 len: 84 ! Aau18262 Novel human DNA-binding protein #1
 (R,K){20,20}
 (K){20}
63: ANAS KKKKKKKKKKKKKKKKKKK XG
 (K){20}
1
AAU18263 ck: 7578 len: 31 ! Aau18263 Novel human DNA-binding protein #1
 (R,K){20,20}
 (K){20}
6: LTEL KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
7: TELK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
8: ELEK KKKKKKKKKKKKKKKKKKK KKKK
 (K){20}
9: LEKK KKKKKKKKKKKKKKKKKKK KKK
 (K){20}
10: EKKK KKKKKKKKKKKKKKKKKKK KK
 (K){20}
11: KKKK KKKKKKKKKKKKKKKKKKK X
 (K){20}
1
AAU18264 ck: 3915 len: 57 ! Aau18264 Novel human DNA-binding protein #1

```
1
(R,K){20,20}
31: KQULL KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
32: QULLK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
33: LLLK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
34: LKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20} XGGF

AAU18265 ck: 3679 len: 37 ! Aau18265 Novel human DNA-binding protein #1
(R,K){20,20}
15: ISPLT KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
16: SPLTK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
17: PLTKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20} X

AAU18266 ck: 657 len: 196 ! Aau18266 Novel human DNA-binding protein #1
(R,K){20,20}
169: FVYFE KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
170: VYFEK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
171: XFEKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
172: FEKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
173: EKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
174: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
175: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
176: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20} X

AAU18267 ck: 4672 len: 57 ! Aau18267 Novel human DNA-binding protein #1
(R,K){20,20}
28: DKTFF KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
29: KTFHK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
30: TFHKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
31: FHKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
32: HKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20} XPGG
```

```
1
AAU18268 ck: 9656 len: 66 ! Aau18268 Novel human DNA-binding protein #1
(R,K){20,20}
38: WVISV KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
39: VISVK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
40: ISVKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
41: SVKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
42: VKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
43: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20} EXKK

AAU18270 ck: 4665 len: 34 ! Aau18270 Novel human DNA-binding protein #1
(R,K){20,20}
10: PELLK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
11: ELLKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
12: LLLKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
13: LKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
14: LKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}

AAU18271 ck: 7810 len: 64 ! Aau18271 Novel human DNA-binding protein #1
(R,K){20,20}
37: LKYFW KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
38: KYFWK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
39: YFWKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
40: FWKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
41: WKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20} XGXP

AAU18272 ck: 269 len: 76 ! Aau18272 Novel human DNA-binding protein #1
(R,K){20,20}
37: TFSRA KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
38: PSRAK KKKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
```

39: SRAXK KKKKKKKKKKKKKKKKKKKKK (K) {20}
40: RAKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
41: AKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
42: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
43: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
44: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}

AAU18273 ck: 8370 len: 45 i Aau18273 Novel human DNA-binding protein #1

17: APTQK KKKKKKKKKKKKKKKKKKKKK (R,K) {20,20}
18: PKTQK KKKKKKKKKKKKKKKKKKKKK (K) {20}
19: KTQKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
20: TQKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
21: QKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
22: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
23: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
24: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
25: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}

AAU18274 ck: 1663 len: 87 i Aau18274 Novel human DNA-binding protein #1

36: KWSXX KKKKKKKKKKKKKKKKKKKKK (R,K) {20,20}
37: WSSXX KKKKKKKKKKKKKKKKKKKKK (K) {20}
38: SSXXK KKKKKKKKKKKKKKKKKKKKK (K) {20}
39: SXKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
40: XKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
41: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}

42: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
43: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
44: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
45: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
46: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
47: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
48: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
49: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
50: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
51: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
52: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
53: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
54: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
55: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
56: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
57: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
58: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
59: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
60: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
61: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
62: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) {20}
63: KKKKK KKKKKKKKKKKKKKKKKKKKK (R,K) {20}

AAU18275 ck: 5607 len: 63 i Aau18275 Novel human DNA-binding protein #1

26: MVELE KKKKKKKKKKKKKKKKKKKKK (R,K) {20,20}
 (K) {20}

27: VLEKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 28: ELEKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 29: LEKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 30: EKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 31: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 32: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 33: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 34: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 35: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 36: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 37: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 38: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 39: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 40: KKKKK KKKKKKKKKKKKKKKKKKK KKKK

AAU18276 ck: 5997 len: 58 i Aau18276 Novel human DNA-binding protein #1
 (R,K) {20,20}
 28: RPTRP KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 29: PTPRP KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}

AAU18277 ck: 5764 len: 74 i Aau18277 Novel human DNA-binding protein #1
 (R,K) {20,20}
 40: EPLSA KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 41: FLSAK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 42: LSAKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 43: SAKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 44: AKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 45: KKKKK KKKKKKKKKKKKKKKKKKK KKKK

46: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 47: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 48: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 49: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 50: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 51: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 52: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 53: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 54: KKKKK KKKKKKKKKKKKKKKKKKK KKKK

AAU18278 ck: 7734 len: 97 i Aau18278 Novel human DNA-binding protein #1
 (R,K) {20,20}
 57: RGCST KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 58: GCSYK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 59: CSYKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 60: SYKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 61: YKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 62: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 63: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 64: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 65: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 66: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 67: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 68: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 69: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 70: KKKKK KKKKKKKKKKKKKKKKKKK KKKK

33: KKKR KKKRRRRRRRRRRRRRRRRRR
(R,K){20}
34: KKKR KKKRRRRRRRRRRRRRRRRRR
(R,K){20}
35: KKKR KKKRRRRRRRRRRRRRRRRRR
(R,K){20}
36: KKKR KKKRRRRRRRRRRRRRRRRRR
(R){20}
37: KKKR KKKRRRRRRRRRRRRRRRRRR
(R){20}
38: KKKR KKKRRRRRRRRRRRRRRRRRR
(R){20}
39: KKKR KKKRRRRRRRRRRRRRRRRRR
(R){20}
40: KKKR KKKRRRRRRRRRRRRRRRRRR
(R){20}
41: KRRR RRRRRRRRRRRRRRRRRRR
(R){20}
42: RRRR RRRRRRRRRRRRRRRRRRR
(R){20}
43: RRRR RRRRRRRRRRRRRRRRRRR
(R){20}
44: RRRR RRRRRRRRRRRRRRRRRRR
(R){20}
45: RRRR RRRRRRRRRRRRRRRRRRR
(R){20}
AAM54700 ck: 1334 len: 86 1 Aam54700 Human brain expressed single exon
1
57: EEEG RRRKKKKKKKKKKKKKKKKKK
(R,K){20,20}
(R,K){20}
58: EEEG RRRKKKKKKKKKKKKKKKKKK
(R,K){20}
59: EEEG RRRKKKKKKKKKKKKKKKKKK
(R,K){20}
60: EEEG RRRKKKKKKKKKKKKKKKKKK
(R,K){20}
61: GRRR KKKKKKKKKKKKKKKKKKK
(R,K){20}
62: RRRR KKKKKKKKKKKKKKKKKKK
(R,K){20}
63: RRRR KKKKKKKKKKKKKKKKKKK
(R,K){20}
64: RRRR KKKKKKKKKKKKKKKKKKK
(R,K){20}
65: KKKR KKKKKKKKKKKKKKKKKKK
(R,K){20}
66: KKKR KKKKKKKKKKKKKKKKKKK
(R,K){20}

67: KKKR KKKKKKKKKKKKKKKKKKK
AAM54795 ck: 9082 len: 167 1 Aam54795 Human brain expressed single exon
1
33: EGGG RRRRRRRRRRRRRRRRRRR
(R,K){20,20}
(R,K){20}
34: EGGG RRRRRRRRRRRRRRRRRRR
(R,K){20}
35: GGGG RRRRRRRRRRRRRRRRRRR
(R,K){20}
AAM55320 ck: 2276 len: 89 1 Aam55320 Human brain expressed single exon
1
23: EEEE KKKKKKKKKKKKKKKKKKK
(R,K){20,20}
(R,K){20}
46: KKEE KKKKKKKKKKKKKKKKKKK
(K){20}
47: KKEE KKKKKKKKKKKKKKKKKKK
(K){20}
48: EEEK KKKKKKKKKKKKKKKKKKK
(R,K){20}
49: EEEK KKKKKKKKKKKKKKKKKKK
(R,K){20}
AAM55623 ck: 1939 len: 130 1 Aam55623 Human brain expressed single exon
1
42: EGGG RRRRRRRRRRRRRRRRRRR
(R,K){20,20}
(R,K){20}
43: GRRR RRRRRRRRRRRRRRRRRRR
(R,K){20}
44: RRRR RRRRRRRRRRRRRRRRRRR
(R,K){20}
45: KERR RRRRRRRRRRRRRRRRRRR
(R,K){20}
46: EERR RRRRRRRRRRRRRRRRRRR
(R,K){20}
47: RRRR RRRRRRRRRRRRRRRRRRR
(R,K){20}
48: RRRR RRRRRRRRRRRRRRRRRRR
(R,K){20}
49: RRRR RRRRRRRRRRRRRRRRRRR
(R,K){20}
50: RRRR RRRRRRRRRRRRRRRRRRR
(R,K){20}
51: RRRR RRRRRRRRRRRRRRRRRRR
(R,K){20}
52: RRRR RRRRRRRRRRRRRRRRRRR
(R,K){20}
53: RRRR RRRRRRRRRRRRRRRRRRR
(R,K){20}

54: RRRR (R,K){20}
RRRRRRRRRRRRRRRRRR EEEEE

AAm55931 ck: 3607 len: 88 i Aam55931 Human brain expressed single exon

1

(R,K){20,20}
(K){20}

39: ERRE KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

40: RRER KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

41: KREK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

42: REKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

43: EKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

44: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

45: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

46: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

47: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

48: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

49: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

50: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

51: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

52: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

53: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

54: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

55: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

56: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

57: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

58: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

59: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

60: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

61: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

62: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

63: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

64: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

65: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

66: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

67: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

68: KKKK KKKKKKKKKKKKKKKKKKK KKKK

(K){20}

69: KKKK KKKKKKKKKKKKKKKKKKK KKKK

1

(R,K){20,20}

(K){20}

1: KKKKKKKKKKKKKKKKKKK KKKK

2: K KKKKKKKKKKKKKKKKK KKKK

(K){20}

3: KK KKKKKKKKKKKKKKKKK KKKK

(K){20}

4: KKK KKKKKKKKKKKKKKKKK KKKK

(K){20}

5: KKK KKKKKKKKKKKKKKKKK KKKK

(K){20}

6: KKKK KKKKKKKKKKKKKKKKK KKKK

(K){20}

7: KKKK KKKKKKKKKKKKKKKKK KKKK

(K){20}

8: KKKK KKKKKKKKKKKKKKKKK KKKK

(K){20}

9: KKKK KKKKKKKKKKKKKKKKK KKKK

(K){20}

10: KKKK KKKKKKKKKKKKKKKKK KKKK

(K){20}

11: KKKK KKKKKKKKKKKKKKKKK EEEE

AAm58224 ck: 1560 len: 88 i Aam58224 Human brain expressed single exon i

1

(R,K){20,20}

(R){20}

43: RRRG RRRRRRRRRRRRRRRRR RRRR

(R){20}

[illegible]

1

7: TEREK KKKRKKKKKKKKKKKKKK KKKK (R,K){20}

8: EREKK KKKKKKKKKKKKKKKKK KKKK (R,K){20}

9: REKKK KKKKKKKKKKKKKKKKK KKKK (R,K){20}

AAM6353 ck: 2394 len: 57 ! Aam6353 Human brain expressed single exon

20: EEEEG RRRRRRRRRRRRRRRRR RRRR (R,K){20,20}

21: EEEGR RRRRRRRRRRRRRRRRR RRRG (R){20}

22: EEGR RRRRRRRRRRRRRRRRR RRRG (R){20}

23: EGRR RRRRRRRRRRRRRRRRR RRRG (R){20}

24: GRRR RRRRRRRRRRRRRRRRR RRRR (R){20}

25: RRRR RRRRRRRRRRRRRRRRR RRRR (R){20}

AAM64090 ck: 4228 len: 24 ! Aam64090 Human brain expressed single exon

1: RRRRRKKKKRRRRRRRRRR RRRR (R,K){20,20}

2: R RRRRRKKKKRRRRRRRRRR RRRR (R,K){20}

3: RR RRRRRKKKKRRRRRRRRRR RRRR (R,K){20}

4: RRR RRRRRKKKKRRRRRRRRRR RRRR (R,K){20}

1

12: KKKKN KKKKKKKKKKKKKKKKK KKKK (R,K){20,20}

13: KKKNK KKKKKKKKKKKKKKKKK KKKK (R,K){20}

14: KKNKK KKKKKKKKKKKKKKKKK KKKK (R,K){20}

15: KNKKK KKKKKKKKKKKKKKKKK KKKK (R,K){20}

16: NKKKK KKKKKKKKKKKKKKKKK KKKK (R,K){20}

17: KKKKK KKKKKKKKKKKKKKKKK KKKK (R,K){20}

18: KKKKK KKKKKKKKKKKKKKKKK KKKK (R,K){20}

AAM64863 ck: 3301 len: 52 ! Aam64863 Human brain expressed single exon

19: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
20: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
21: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
22: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
23: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
24: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
25: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
26: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
27: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
28: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
29: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
30: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
31: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

AAm65355 ck: 4895 len: 51 ! Aam65355 Human brain expressed single exon
(R,K){20,20}
(R,K){20}
18: LFKPM RKKKKKKKKKKKKKKKKKK KCLTT
(R,K){20}
19: FKPMR KKKKKKKKKKKKKKKKKKK KCLTT
(R,K){20}
20: KPMRK RKKKKKKKKKKKKKKKKKK LTTT
(R,K){20}

AAm66249 ck: 5383 len: 86 ! Aam66249 Human bone marrow expressed probe
(R,K){20,20}
(R,K){20}
15: RRRRG RRRKKKKKKKKKKKKKKKK KKKRR
(R,K){20}
16: RRRGR RRRKKKKKKKKKKKKKKKK KRRRR
(R,K){20}
17: RRGRR RRRKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
18: RGRRR RRRKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
19: GRRRR RKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
20: RRRRR KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}

(R,K){20}
21: RRRRK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
22: RRRKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
23: RRRKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
24: RKKKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
25: KKKKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
26: KKKKK RKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
27: KKKKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
28: KKKKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
29: KKKKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
30: KKKKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
31: RKKKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
32: KKKKK RKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
33: KKKKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
34: KKKKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
35: KKKKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
36: KKKKK KKKKKKKKKKKKKKKKKKK RRRRR
(R,K){20}
37: RKKKK RRRRRKKKKKKKKKKKKKK RRRRR
(R){20}
38: KKKKK RRRRRKKKKKKKKKKKKKK RRRRR
(R){20}
39: KKKKK RRRRRKKKKKKKKKKKKKK RRRRR
(R){20}
40: KKKKK RRRRRKKKKKKKKKKKKKK RRRRR
(R){20}
41: KRRRR RRRRRKKKKKKKKKKKKKK RRRRR
(R){20}
42: RRRRR RRRRRKKKKKKKKKKKKKK RRRRR
(R){20}
43: RRRRR RRRRRKKKKKKKKKKKKKK RRRRR
(R){20}
44: RRRRR RRRRRKKKKKKKKKKKKKK RRRRR
(R){20}

```

45: RRRR RRRRRRRRRRRRRRRR NKQTK
AAM67100 ck: 1334 len: 86 ! Aam67100 Human bone marrow expressed probe
1
57: EEEEG RRRRRRRRRRRRRRRR KKKKK
(R,K){20,20}
(R,K){20}
58: EEEGR RRRRRRRRRRRRRRRR KKKKK
(R,K){20}
59: EEEGR RRRRRRRRRRRRRRRR KKKKK
(R,K){20}
60: EGRRR RRRRRRRRRRRRRRRR KKKKK
(R,K){20}
61: GRRRR RRRRRRRRRRRRRRRR KKKKK
(R,K){20}
62: RRRRR RRRRRRRRRRRRRRRR KKKKK
(R,K){20}
63: RRRRR RRRRRRRRRRRRRRRR KKKKK
(R,K){20}
64: RRRRR RRRRRRRRRRRRRRRR KKK
(R,K){20}
65: RRRRR RRRRRRRRRRRRRRRR KK
(R,K){20}
66: RRRRR RRRRRRRRRRRRRRRR K
(R,K){20}
67: RRRRR RRRRRRRRRRRRRRRR KKKKK
(R,K){20}
AAM67180 ck: 9082 len: 167 ! Aam67180 Human bone marrow expressed probe
1
33: EEEGR RRRRRRRRRRRRRRRR RRRGG
(R,K){20,20}
(R,K){20}
34: EEEGR RRRRRRRRRRRRRRRR RRRGG
(R,K){20}
35: EEEGR RRRRRRRRRRRRRRRR RRRGG
(R,K){20}
AAM6717 ck: 2276 len: 89 ! Aam6717 Human bone marrow expressed probe
1
23: EEEEG RRRRRRRRRRRRRRRR EEEKK
(R,K){20,20}
(R,K){20}
46: EEEEG RRRRRRRRRRRRRRRR KRRKE
(R,K){20}
47: EEEGR RRRRRRRRRRRRRRRR RRRKE
(R,K){20}
48: EEEGR RRRRRRRRRRRRRRRR RRRKE
(R,K){20}
49: EEEGR RRRRRRRRRRRRRRRR RRRKE
(R,K){20}
AAM68007 ck: 1939 len: 130 ! Aam68007 Human bone marrow expressed probe

```

```

1
42: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20,20}
(R,K){20}
43: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
44: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
45: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
46: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
47: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
48: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
49: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
50: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
51: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
52: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
53: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
54: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
AAM68298 ck: 3607 len: 88 ! Aam68298 Human bone marrow expressed probe
1
39: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20,20}
(R,K){20}
40: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
41: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
42: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
43: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
44: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
45: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
46: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}
47: EEEGR RRRRRRRRRRRRRRRR RRRRR
(R,K){20}

```

48: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 49: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 50: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 51: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 52: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 53: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 54: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 55: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 56: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 57: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 58: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 59: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 60: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 61: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 62: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 63: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 64: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 65: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 66: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 67: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 68: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 69: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}

AA6869 ck: 3937 len: 85 i AA6869 Human bone marrow expressed probe

1
 1: KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}

2: K KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 3: K KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 4: K KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 5: K KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 6: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 7: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 8: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 9: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 10: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 11: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}

AA69569 ck: 2686 len: 71 i AA69569 Human bone marrow expressed probe

1
 20: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20, 20}
 (K) {20}
 21: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 22: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 23: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 24: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 25: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20}
 26: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20}
 27: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20}
 28: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20}
 29: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20}
 30: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20}
 31: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20}
 32: KKKK KKKKKKKKKKKKKKKKKKK KKKK
 (R, K) {20}


```

33: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
34: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
35: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
36: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
37: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
38: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
39: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
40: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
41: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
42: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
43: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
44: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
45: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
46: KKKKK KKKKKKKKKKKKKKKKKKK KKKSA
    (R,K) {20}
47: KKKKK KKKKKKKKKKKKKKKKKKK KKSAA
    (K) {20}
48: KKKKK KKKKKKKKKKKKKKKKKKK KSAA
    (K) {20}
49: KKKKK KKKKKKKKKKKKKKKKKKK SAA
    (K) {20}

AAM70678 ck: 1560 len: 88 ! Aam70678 Human bone marrow expressed probe
    (R,K) {20,20}
43: RRRRG RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
44: RRRGR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
45: RRGRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
46: RGRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
47: GRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
48: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}

```

```

49: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
50: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
51: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
52: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
53: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
54: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
55: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
56: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
57: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
58: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
59: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
60: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
61: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}
62: RRRRR RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}

AAM70881 ck: 2324 len: 36 ! Aam70881 Human bone marrow expressed probe
    (R,K) {20,20}
16: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
17: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}

AAM7367 ck: 8343 len: 66 ! Aam7367 Human bone marrow expressed probe
    (R,K) {20,20}
6: ETERE KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
7: TEREK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
8: EREKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}
9: REKKK KKKKKKKKKKKKKKKKKKK KKKKK
    (R,K) {20}

AAM76347 ck: 2394 len: 57 ! Aam76347 Human bone marrow expressed probe
    (R,K) {20,20}
20: EEEEG RRRRRRRRRRRRRRRRRR RRRR
    (R) {20}

```

21: EEEGR RRRRRRRRRRRRRRRRRRRRRRRRRG
(R) {20}

22: EEEGR RRRRRRRRRRRRRRRRRRRRRRRRRG
(R) {20}

23: EEEGR RRRRRRRRRRRRRRRRRRRRRRRRRG
(R) {20}

24: GRRRR RRRRRRRRRRRRRRRRRRRRRRRRRG
(R) {20}

25: RRRRR RRRRRRRRRRRRRRRRRRRRRRRRRG
(R) {20}

AAM76911 ck: 4228 len: 24 ! Aam76911 Human bone marrow expressed probe

```
1      (R,K){20,20}
      (R,K){20}
1:    RRRRRRRKKKKRRRRRRRRR RKR
```

```
2: R RRRRRKKKKRRRRRRRRRR KRT
```

```

3:      RR RRRRRKKKKRRRRRRRRRRRK RT
          (R,K){20}

```

```

4:      RRR RRRKKKKRRRRRRRRRRR T
      (R,K){20}

```

AAM78048 ck: 4895 len: 51 | Aam78048 Human bone marrow expressed probe

```

1      (R,K){20,20}
      (R,K){20}
18: LFKPM RRRRRKKRRKKRRKKRR KLLTT

```

19: $(R, K) \{20\}$
 KRRKKRRKKKKRRK KLTTT

20: KPMRK RRRKKRRKKKKRRKK (R, K) { 20 } LTTT

AAM82533 ck: 1736 len: 40 ! Aam82533 Human immune/haematopoietic antigen

$$\begin{array}{c} (R, K) \{20, 20\} \\ (K) \{20\} \end{array}$$

(K) {20}

```
(K) { 20 }
20: GSIKK KKKKKKKKKKKKKKKKKKKKK
```

(K) {20}

21: SLKK KKKKKKKKKKKKKKKKK

AAm85748 ck: 7503 Len: 74 ! Aam85748 Human immune/haematopoietic antigen

$$(R, K) \begin{cases} 20, 20 \\ (K) \end{cases} \{20\}$$

```

46: HLTDK KKKKKKKKKKKKKKKKKKKKK (K) { 20 } KKKKK

```

(K) { 20 }

47: LTKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

48: TDIKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) { 20 }

49 : DKKK KKKKKKKKKKKKKKKKK KKKK

(K) { 20 }

50 : KKKKK KKKKKKKKKKKKKKKKK KKKKK

51: KKKKK KKKKKKKKKKKKKKKKKKK KKKK (K) { 20 }

52: KKKKK KKKKKKKKKKKKKKKKKKK KKK (K) {20}

(K) { 20 }

53 : KKKKK KKKKKKKKKKKKKKKKK KK

54: KKKKK KKKKKKKKKKKKKKKKKKK K (K) {20}

(K) { 20 }

55 : KKKKK KKKKKKKKKKKKKKKKKKK

AAM90546 ck: 6676 len: 74 | Aam90546 Human immune/haematopoietic antigen

```

1      (R,K) { 20, 20 }
      (K) { 20 }
47. 1BTE0 KKKKKKKKKKKKKKKKKKKKK KKKKK

```

```
(K) { 20 }
48: RTFOK KKKKKKKKKKKKKKKKKKKKK KKKKK
```

```

      (K) { 20 }
49: TFOKK KKKKKKKKKKKKKKKKKKK KKKX

```

```

50: FOKKK KKKKKKKKKKKKKKKKKKK KXXG
      (K) { 20 }

```

51: QKKKK KKKKKKKKKKKKKKKKK KXXC
(K) { 20 }

52: KKKKK KKKKKKKKKKKKKKKKKKKKK (K) { 20 } XXG

AAM90618 ck: 5691 len: 108 ! Aam90618 Human immune/haematopoietic antigen

$$\begin{array}{l} 1 \\ (R, K) \{20, 20\} \\ (K) \{20\} \end{array}$$
$$(K) \{20\}$$

(K) {20}

$$(K) \{20\}$$

(K) {20}

(K) {20}

83 . KKKKKKKKKKKKKKKKKKKKK

(K) {20}

 $(K) \{20\}$

85: KKKKK KKKKKKKKKKKKKKKKKKK KKKK
 (K) {20}
 86: KKKKK KKKKKKKKKKKKKKKKKKK KKK
 (K) {20}
 87: KKKKK KKKKKKKKKKKKKKKKKKK KK
 (K) {20}
 88: KKKKK KKKKKKKKKKKKKKKKKKK K
 (K) {20}
 89: KKKKK KKKKKKKKKKKKKKKKKKK

1
 AAm91030 ck: 8102 len: 111 ! Aam91030 Human immune/haematopoietic antigen
 (R,K) {20,20}
 (K) {20}

78: EFHIL KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 79: PHILK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 80: HILKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 81: ILKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 82: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 83: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 84: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 85: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 86: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 87: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 88: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 89: KKKKK KKKKKKKKKKKKKKKKKKK KKK
 (K) {20}
 90: KKKKK KKKKKKKKKKKKKKKKKKK KK
 (K) {20}
 91: KKKKK KKKKKKKKKKKKKKKKKKK K
 (K) {20}
 92: KKKKK KKKKKKKKKKKKKKKKKKK

1
 AAm91162 ck: 1109 len: 98 ! Aam91162 Human immune/haematopoietic antigen
 (R,K) {20,20}
 (K) {20}
 53: QTKNT KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 54: TQNTK KKKKKKKKKKKKKKKKKKK KKKKK

1

55: KNTKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 56: NTKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 57: TKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 58: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 59: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 60: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 61: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 62: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 63: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 64: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 65: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 66: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 67: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 68: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}

1
 AAm91891 ck: 8102 len: 111 ! Aam91891 Human immune/haematopoietic antigen
 (R,K) {20,20}
 (K) {20}
 78: EFHIL KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 79: PHILK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 80: HILKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 81: ILKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 82: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 83: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 84: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 85: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
 (K) {20}
 86: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

87: KKKKK (K) {20}
88: KKKKK (K) {20}
89: KKKKK (K) {20}
90: KKKKK (K) {20}
91: KKKKK (K) {20}
92: KKKKK (K) {20}

1 AAM91933 ck: 8285 len: 55 ! Aam91933 Human digestive system antigen SEQ

(R,K) {20,20}

33: PPTRP KKKKKKKKKKKKKKKKKKK KKK

(K) {20}

34: PTPPK KKKKKKKKKKKKKKKKKKK GK

1 AAM92433 ck: 1663 len: 87 ! Aam92433 Human digestive system antigen SEQ

(R,K) {20,20}

(K) {20}

36: KMSXK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

37: WSSXK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

38: SSXKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

39: SXKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

40: XKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

41: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

42: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

43: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

44: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

45: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

46: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

47: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

48: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

49: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

50: KKKKK (K) {20}

(K) {20}

51: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

52: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

53: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

54: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

55: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

56: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

57: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

58: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

59: KKKKK KKKKKKKKKKKKKKKKKKK KKKRG

(K) {20}

60: KKKKK KKKKKKKKKKKKKKKKKKK KKKGX

(K) {20}

61: KKKKK KKKKKKKKKKKKKKKKKKK KRGXP

(K) {20}

62: KKKKK KKKKKKKKKKKKKKKKKKK RGXPF

(R,K) {20}

63: KKKKK KKKKKKKKKKKKKKKKKKK GXPFK

1 AAO00092 ck: 9065 len: 113 ! Aao00092 Human polypeptide SEQ ID NO 13984.

(R,K) {20,20}

(K) {20}

91: CLGCL KKKKKKKKKKKKKKKKKKK KPF

(K) {20}

92: LGCLK KKKKKKKKKKKKKKKKKKK FPF

1 AAO00222 ck: 2916 len: 132 ! Aao00222 Human polypeptide SEQ ID NO 14114.

(R,K) {20,20}

(K) {20}

28: XPLPP KKKKKKKKKKKKKKKKKKK GPPPK

1 AAO00232 ck: 1000 len: 102 ! Aao00232 Human polypeptide SEQ ID NO 14124.

(R,K) {20,20}

(K) {20}

29: RDCFP KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

30: DCFPK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

31: CFFKK KKKKKKKKKKKKKKKKKKK KKKKK

[illegible]

```

1      AAO00291 ck: 8100 len: 124   ! Aao00291 Human polypeptide SEQ ID NO 14183.
          (R,K){20,20}
          (K){20}
22: QHFCM KKKKKKKKKKKKKKKKKKK FFKKG

1      AAO00439 ck: 6396 len: 122   ! Aao00439 Human polypeptide SEQ ID NO 14331.
          (R,K){20,20}
          (K){20}
23: CLMLV KKKKKKKKKKKKKKKKKKK KKKKK
24: LMLVK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
25: WLVRK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
26: LVRRK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
27: VKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
28: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
29: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
30: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
31: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
32: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
33: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
34: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
35: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
36: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
37: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
38: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
39: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
40: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
41: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (K){20}
42: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (R,K){20}
43: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
          (R,K){20}

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51: KKKK (K) {20} KKKK
52: KKKK (K) {20} KKKK
53: KKKK (K) {20} KQ
54: KKKK (K) {20} Q
AA002186 ck: 4844 len: 57 ! Aa002186 Human polypeptide SEQ ID NO 16078.
(R, K) {20, 20}
11: HCCL KKKKKKKKKKKKKKKKKKKKK KKKK
12: CCLL (K) {20} KKKK
AA002310 ck: 704 len: 137 ! Aa002310 Human polypeptide SEQ ID NO 16202.
(R, K) {20, 20}
22: HSLN KKKKKKKKKKKKKKKKKKKR GGSVK
1
AA002477 ck: 2018 len: 112 ! Aa002477 Human polypeptide SEQ ID NO 16369.
(R, K) {20, 20}
83: LASAV KKKKKKKKKKKKKKKKKKK KKKK
84: ASAV (K) {20} KKKK
85: SAVK (K) {20} KKKK
86: AVKK (K) {20} KKKK
87: VKKK (K) {20} KKKK
88: KKKK (K) {20} KKKK
89: KKKK (K) {20} KKKK
90: KKKK (K) {20} KKKK
91: KKKK (K) {20} KKKK
92: KKKK (K) {20} K
93: KKKK (K) {20} KKKK
AA002733 ck: 2860 len: 132 ! Aa002733 Human polypeptide SEQ ID NO 16625.
(R, K) {20, 20}
86: FFPSS KKKKKKKKKKKKKKKKKKK KKKK

1
87: FFSK (K) {20} KKKK
88: FSLK (K) {20} KKKK
89: SLKK (K) {20} KKKK
90: LKKK (K) {20} KKKK
91: KKKK (K) {20} KKKK
92: KKKK (K) {20} KKKK
93: KKKK (K) {20} KKKK
94: KKKK (K) {20} KKKK
95: KKKK (K) {20} KKKK
96: KKKK (K) {20} KKKK
97: KKKK (K) {20} KKKK
98: KKKK (K) {20} KKKK
99: KKKK (K) {20} KKKK
100: KKKK (K) {20} KKKK
101: KKKK (R, K) {20} KKKK
AA002946 ck: 3649 len: 126 ! Aa002946 Human polypeptide SEQ ID NO 16838.
(R, K) {20, 20}
25: DEATS KKKKKKKKKKKKKKKKKKK KKKK
26: EATSK (K) {20} KKKK
27: ATSK (K) {20} KKKK
28: TSKK (K) {20} KKKK
29: SKKK (K) {20} KKKK
30: KKKK (K) {20} KKKK
31: KKKK (K) {20} KKKK
32: KKKK (K) {20} KKKK

(K) {20}
33: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
34: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
35: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
36: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
37: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
38: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

AA002961 ck: 4320 len: 83 i Aa002961 Human polypeptide SEQ ID NO 16853.
(R, K) {20, 20}
22: RMFSS KKKKKKKKKKKKKKKKKKK KTAIT
(K) {20}
23: MFSSK KKKKKKKKKKKKKKKKKKK TAITK

AA003006 ck: 5212 len: 102 i Aa003006 Human polypeptide SEQ ID NO 16898.
(R, K) {20, 20}
59: HDEFP KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
60: DPEPK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
61: FPEPK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
62: FPKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
63: PKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
64: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
65: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
66: KKKKK KKKKKKKKKKKKKKKKKKK KASSS
(K) {20}
67: KKKKK KKKKKKKKKKKKKKKKKKK ASSSS
AA003024 ck: 6865 len: 64 i Aa003024 Human polypeptide SEQ ID NO 16916.
(R, K) {20, 20}
15: SPAKA KKKKKKKKKKKKKKKKKKK KRGGP
(R, K) {20}
16: FAKAR KKKKKKKKKKKKKKKKKKK RGGP
(R, K) {20}
17: AKARK KKKKKKKKKKKKKKKKKKK GGPP

AA003113 ck: 5587 len: 60 i Aa003113 Human polypeptide SEQ ID NO 17005.
(R, K) {20, 20}
33: EIMNL KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
34: INMLK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
35: NMLKK KKKKKKKKKKKKKKKKKKK KRGGG
(K) {20}
36: MLKKK KKKKKKKKKKKKKKKKKKK RGGGL
(R, K) {20}
37: LKKKK KKKKKKKKKKKKKKKKKKK GGGL

AA003123 ck: 6627 len: 108 i Aa003123 Human polypeptide SEQ ID NO 17015.
(R, K) {20, 20}
33: RTWRX KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
34: TWRLX KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
35: WRXXK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
36: RXXKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
37: XKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
38: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
39: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
40: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
41: KKKKK KKKKKKKKKKKKKKKKKKK KRGGG
(K) {20}
42: KKKKK KKKKKKKKKKKKKKKKKKK RGGGQ
(R, K) {20}
43: KKKKK KKKKKKKKKKKKKKKKKKK GGQKK

AA003132 ck: 3903 len: 116 i Aa003132 Human polypeptide SEQ ID NO 17024.
(R, K) {20, 20}
1: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKK KRGGG
(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKK KRGGP
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKK RGGPP

(R,K){20}
6: KKKKK KKKKKKKKKKKKKKKKKR GGPPK

1 AAO03152 ck: 7891 len: 35 i Aao03152 Human polypeptide SEQ ID NO 17044.
(R,K){20,20}
(R,K){20}
13: LIXYT RKKKKKKKKKKKKKKKKK RGG
(R,K){20}
14: IXYTR KKKKKKKKKKKKKKKKKR GG

AAO03168 ck: 7233 len: 67 i Aao03168 Human polypeptide SEQ ID NO 17060.
(R,K){20,20}
(K){20}
17: ELMP KKKKKKKKKKKKKKKKKK KRGA
(K){20}
18: LMPK KKKKKKKKKKKKKKKKKK RGGAP
(R,K){20}
19: LMPK KKKKKKKKKKKKKKKKKR CGAPF

1 AAO03243 ck: 411 len: 95 i Aao03243 Human polypeptide SEQ ID NO 17135.
(R,K){20,20}
(K){20}
51: LHAV KKKKKKKKKKKKKKKKKK KEKK
(K){20}
52: LHAV KKKKKKKKKKKKKKKKKK KEKKR
(K){20}
53: HAVK KKKKKKKKKKKKKKKKKK EKKRG

AAO03273 ck: 2981 len: 88 i Aao03273 Human polypeptide SEQ ID NO 17165.
(R,K){20,20}
(K){20}
43: IXYL KKKKKKKKKKKKKKKKKK KKKK
(K){20}
44: XYLL KKKKKKKKKKKKKKKKKK KKKK
(K){20}
45: YLLK KKKKKKKKKKKKKKKKKK KKKK
(K){20}
46: LLKK KKKKKKKKKKKKKKKKKK KKKK
(K){20}
47: LKKK KKKKKKKKKKKKKKKKKK KKKK
(K){20}
48: KKKK KKKKKKKKKKKKKKKKKK KKKK
(K){20}
49: KKKK KKKKKKKKKKKKKKKKKK KKKK
(K){20}
50: KKKK KKKKKKKKKKKKKKKKKK KKKK
(K){20}
51: KKKK KKKKKKKKKKKKKKKKKK KKKK
(K){20}
52: KKKK KKKKKKKKKKKKKKKKKK KKKK

(K){20}
53: KKKK KKKKKKKKKKKKKKKKKK KKKK
(K){20}
54: KKKK KKKKKKKKKKKKKKKKKK KKPT
(K){20}
55: KKKK KKKKKKKKKKKKKKKKKK KKPTP
(K){20}
56: KKKK KKKKKKKKKKKKKKKKK KPTPK
(K){20}
57: KKKK KKKKKKKKKKKKKKKKK PTPK

1 AAO03277 ck: 2981 len: 89 i Aao03277 Human polypeptide SEQ ID NO 17169.
(R,K){20,20}
(K){20}
1: KKKKKKKKKKKKKKKKKK KKRM
(K){20}
2: K KKKKKKKKKKKKKKKKK KKRM
(K){20}
3: KK KKKKKKKKKKKKKKKKK KRKN
(K){20}
4: KK KKKKKKKKKKKKKKKKK RKNK
(R,K){20}
5: KKK KKKKKKKKKKKKKKKR MNKK

1 AAO03278 ck: 6777 len: 74 i Aao03278 Human polypeptide SEQ ID NO 17170.
(R,K){20,20}
(K){20}
47: RAVQ KKKKKKKKKKKKKKKKK KKGG
(K){20}
48: AVAQ KKKKKKKKKKKKKKKKK KGGG
(K){20}
49: VAQK KKKKKKKKKKKKKKKKK KGGV
(K){20}
50: AQK KKKKKKKKKKKKKKKKK GGVL

1 AAO03284 ck: 9650 len: 115 i Aao03284 Human polypeptide SEQ ID NO 17176.
(R,K){20,20}
(K){20}
22: KALV KKKKKKKKKKKKKKKKK KKKK
(K){20}
23: ALVK KKKKKKKKKKKKKKKKK KKKK
(K){20}
24: LVSK KKKKKKKKKKKKKKKKK KKKR
(K){20}
25: VSKK KKKKKKKKKKKKKKKKK KKRK
(K){20}
26: SKKK KKKKKKKKKKKKKKKKK KRRG
(K){20}
27: KKKK KKKKKKKKKKKKKKKKK KRGA

78: PSRAK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX
79: SPRAK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX
80: PAKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX
81: AKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX
82: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX
83: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX
84: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX
85: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX
86: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX
87: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX
88: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX

1
AAO03700 ck: 1279 len: 43 1 Aao03700 Human polypeptide SEQ ID NO 17592.
^{(R, K) {20, 20}}
^{(K) {20}}
19: YSQRL KKKKKKKKKKKKKKKKKKKKK GGGPF

1
AAO03703 ck: 7016 len: 113 1 Aao03703 Human polypeptide SEQ ID NO 17595.
^{(R, K) {20, 20}}
^{(K) {20}}
58: IMDAE KKKKKKKKKKKKKKKKKKKKK KKKKK
59: WDAEK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKKK
60: DAEKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKKK
61: AEKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKKK
62: EKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKKK
63: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKKK
64: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KSPGG
65: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX SPGGA

1
AAO03766 ck: 8808 len: 81 1 Aao03766 Human polypeptide SEQ ID NO 17658.
^{(R, K) {20, 20}}

49: TTAC ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKK
50: TTACK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKK
51: TACKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKK
52: ACKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKK
53: CKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKK
54: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKK
55: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKK
56: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KGGGG
57: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX GGGGA

1
AAO03841 ck: 8734 len: 100 1 Aao03841 Human polypeptide SEQ ID NO 17733.
^{(R, K) {20, 20}}
^{(K) {20}}
36: KOHYP KKKKKKKKKKKKKKKKKKKKK KKKKK

37: OHYPK KKKKKKKKKKKKKKKKKKKKK KKKKK
38: HYPKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKKK
39: YPKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKKK
40: PKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKKK
41: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKKK
42: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKRTK
43: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KRTKQ
44: KKKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX RTKKK
45: KKKKK ^{(R, K) {20}}XXXXXXXXXXXXXXXXXXXX TKKKK

1
AAO03906 ck: 4312 len: 100 1 Aao03906 Human polypeptide SEQ ID NO 17798.
^{(R, K) {20, 20}}
^{(K) {20}}
34: NKQNK KKKKKKKKKKKKKKKKKKKKK KKKKK
35: KONKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKKK
36: QNKKK ^{(K) {20}}XXXXXXXXXXXXXXXXXXXX KKKKK

37: NQKKK (K) {20}
 38: QKKKK (K) {20}
 39: KKKKK (K) {20}
 40: KKKKK (K) {20}
 41: KKKKK (K) {20}
 42: KKKKK (K) {20}
 43: KKKKK (K) {20}
 44: KKKKK (K) {20}
 45: KKKKK (K) {20}
 46: KKKKK (K) {20}

AA003967 ck: 7917 len: 53 ! Aa003967 Human polypeptide SEQ ID NO 17859.

16: KLKKS (R,K) {20,20}
 17: LKSKK (K) {20}
 18: XKSKK (K) {20}
 19: KSKKK (K) {20}
 20: SKKKK (K) {20}
 21: KKKKK (K) {20}
 22: KKKKK (K) {20}
 23: KKKKK (K) {20}
 24: KKKKK (K) {20}
 25: KKKKK (K) {20}

AA004512 ck: 4277 len: 62 ! Aa004512 Human polypeptide SEQ ID NO 18404.

24: KKKKK (R,K) {20,20}

AA004619 ck: 8654 len: 39 ! Aa004619 Human polypeptide SEQ ID NO 18511.
 13: LITFL (R,K) {20,20}
 14: ITFLK (K) {20}
 15: TFLKK (K) {20}
 16: FLKKK (K) {20}
 17: LKKKK (K) {20}
 18: KKKKK (K) {20}
 19: KKKKK (K) {20}

AA004644 ck: 2038 len: 70 ! Aa004644 Human polypeptide SEQ ID NO 18536.

23: AKPPT (R,K) {20,20}

AA004645 ck: 7825 len: 118 ! Aa004645 Human polypeptide SEQ ID NO 18537.

46: IKSFL (R,K) {20,20}
 47: KSFLR (K) {20}
 48: SFLRK (K) {20}
 49: FLRKK (K) {20}
 50: LRKKK (K) {20}
 51: RKKKK (K) {20}
 52: KKKKK (K) {20}
 53: KKKKK (K) {20}
 54: KKKKK (K) {20}
 55: KKKKK (K) {20}
 56: KKKKK (K) {20}
 57: KKKKK (K) {20}
 58: KKKKK (K) {20}

59: KKKKK (K) {20} KKKR
60: KKKKK (K) {20} KPRG
61: KKKKK (K) {20} KPRG
62: KKKKK (K) {20} PRGG

AA004647 ck: 4805 len: 58 ! Aa004647 Human polypeptide SEQ ID NO 18539.

(R,K) {20,20}
28: KPTRP KKKKK (K) {20} KKKK
29: PTPPK KKKKK (K) {20} KKKK
30: TRPKK KKKKK (K) {20} KKKK
31: RPKKK KKKKK (K) {20} KKKK
32: PKKKK KKKKK (K) {20} KKKK
33: KKKKK KKKKK (K) {20} KKKK
34: KKKKK KKKKK (K) {20} KKKK
35: KKKKK KKKKK (K) {20} KKKK
36: KKKKK KKKKK (K) {20} KKS
37: KKKKK KKKKK (K) {20} KS
38: KKKKK KKKKK (K) {20} S

AA004674 ck: 2036 len: 33 ! Aa004674 Human polypeptide SEQ ID NO 18566.

(R,K) {20,20}
9: WCYIT KKKKK (K) {20} KARG
10: CYITK KKKKK (K) {20} KARG
11: YITKK KKKKK (K) {20} ARG

AA004679 ck: 7190 len: 31 ! Aa004679 Human polypeptide SEQ ID NO 18571.

(R,K) {20,20}
8: CWFTQ KKKKK (K) {20} KDRG
9: WFTQK KKKKK (K) {20} DRG

AA004682 ck: 5605 len: 60 ! Aa004682 Human polypeptide SEQ ID NO 18574.

(R,K) {20,20}
2: L KKKKK (K) {20} KKKK
3: LK KKKKK (K) {20} KKQD
4: LKK KKKKK (K) {20} KQDL
5: LKKK KKKKK (K) {20} QKDL

AA004690 ck: 7157 len: 81 ! Aa004690 Human polypeptide SEQ ID NO 18582.

(R,K) {20,20}
1: KKKKK (K) {20} KKKK
2: K KKKKK (K) {20} KKKK
3: KK KKKKK (K) {20} KKKK
4: KKK KKKKK (K) {20} KKKK
5: KKKK KKKKK (K) {20} KKKK
6: KKKKK KKKKK (K) {20} KKKK
7: KKKKK KKKKK (K) {20} KKKK
8: KKKKK KKKKK (K) {20} PKKK

AA004715 ck: 6984 len: 35 ! Aa004715 Human polypeptide SEQ ID NO 18607.

(R,K) {20,20}
8: LGSKD KKKKK (K) {20} KKKK
9: GSKDK KKKKK (K) {20} KKKK
10: SKDKK KKKKK (K) {20} AKKD

AA004743 ck: 9412 len: 54 ! Aa004743 Human polypeptide SEQ ID NO 18635.

(R,K) {20,20}
21: KINKL KKKKK (K) {20} KRAA
22: INKLK KKKKK (K) {20} RAAR
23: NKLKK KKKKK (R,K) {20} AAAR

AA004747 ck: 8399 len: 39 ! Aa004747 Human polypeptide SEQ ID NO 18639.

1

(R,K){20,20}

(K){20}

13: RTGFV KKKKKKKKKKKKKKKKKKK KKRGG

(K){20}

14: TGFVK KKKKKKKKKKKKKKKKKKK KRGGG

(K){20}

15: GFVKK KKKKKKKKKKKKKKKKKKK RGGGF

(R,K){20}

16: FVKKK KKKKKKKKKKKKKKKKKKK RGGGF

AA004752 ck: 1021 len: 71 ! Aa004752 Human polypeptide SEQ ID NO 18644.

(R,K){20,20}

(K){20}

19: QEOGL KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

20: EOGLK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

21: OGLKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

22: GLKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

23: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

24: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

25: KKKKK KKKKKKKKKKKKKKKKKKK KKGKG

(K){20}

26: KKKKK KKKKKKKKKKKKKKKKKKK KGGGL

(K){20}

27: KKKKK KKKKKKKKKKKKKKKKKKK GGGLL

AA004755 ck: 5521 len: 59 ! Aa004755 Human polypeptide SEQ ID NO 18647.

(R,K){20,20}

(K){20}

31: INSLE KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

32: NSLEK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

33: SLEKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

34: LEKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

35: EKKKK KKKKKKKKKKKKKKKKKKK KKKKT

(K){20}

36: KKKKK KKKKKKKKKKKKKKKKKKK KKKTT

(K){20}

37: KKKKK KKKKKKKKKKKKKKKKKKK KKT

(K){20}

38: KKKKK KKKKKKKKKKKKKKKKKKK KT

(K){20}

39: KKKKK KKKKKKKKKKKKKKKKKKK T

AA004756 ck: 6952 len: 26 ! Aa004756 Human polypeptide SEQ ID NO 18648.

(R,K){20,20}

(K){20}

4: FFY KKKKKKKKKKKKKKKKKKK SSS

AA004758 ck: 7836 len: 115 ! Aa004758 Human polypeptide SEQ ID NO 18650.

(R,K){20,20}

(K){20}

7: PFYQL KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

8: FYQLK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

9: YQLKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

10: QLKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

11: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

12: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

13: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

14: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

15: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

16: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

17: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

18: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

19: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

20: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

21: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

22: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

23: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

24: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

25: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

26: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

[illegible]

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1.
AA004764 ck: 2035 len: 54 ! Aa004764 Human polypeptide SRQ ID NO 18656.
(R,K){20,20}
(K){20}
9: GDSL KKKKKKKKKKKKKKKKKKKKKKKKK
10: DSLK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
11: SLKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
12: SLKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
13: LKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
14: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}

AA004802 ck: 1223 len: 105 ! Aa004802 Human polypeptide SRQ ID NO 18694.
(R,K){20,20}
(K){20}
34: FFSRQ KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}

AA004854 ck: 4121 len: 58 ! Aa004854 Human polypeptide SRQ ID NO 18746.
(R,K){20,20}
(K){20}
13: TPFRA KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
14: PFRK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
15: FRKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
16: RAKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
17: AKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
18: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
19: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
20: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
21: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
22: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
23: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
24: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
25: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}
26: KKKK KKKKKKKKKKKKKKKKKKKKKKKK
(K){20}

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27: KKKKK (K) {20}
28: KKKKK (K) {20}
29: KKKKK (K) {20}
30: KKKKK (K) {20}
31: KKKKK (K) {20}
32: KKKKK (R,K) {20} GGNFK

AA004856 ck: 4861 len: 135 ! Aa004856 Human polypeptide SEQ ID NO 18748.

41: LTLTT (R,K) {20,20}
42: TLTTK (K) {20}
43: LTTKK (K) {20}
44: TTYKK (K) {20}
45: TTKKK (K) {20}
46: KKKKK (K) {20}
47: KKKKK (K) {20}
48: KKKKK (K) {20}
49: KKKKK (K) {20}
50: KKKKK (K) {20}
51: KKKKK (K) {20}
52: KKKKK (K) {20}
53: KKKKK (K) {20} GGGGP
AA004872 ck: 3224 len: 93 ! Aa004872 Human polypeptide SEQ ID NO 18764.
(R,K) {20,20}
8: FLVYL KKKKKKKKKKKKKKKKKKK KKKRG
9: LVYLL KKKKKKKKKKKKKKKKKKK KRGKG (K) {20}

10: YKLKK KKKKKKKKKKKKKKKKKKK KRGSP
11: KLKKK KKKKKKKKKKKKKKKKKKK RGGPL
12: LKKKK (R,K) {20} GGPLK

AA004873 ck: 7719 len: 66 ! Aa004873 Human polypeptide SEQ ID NO 18765.

21: SFLIE (R,K) {20,20} KGGGP
22: FLIEK (K) {20} GGGPL

AA004874 ck: 5753 len: 58 ! Aa004874 Human polypeptide SEQ ID NO 18766.

10: SEKLP (R,K) {20,20} KKKKK
11: EKLPR (K) {20} KKKKI
12: KLPKK (K) {20} GKLIK

AA004881 ck: 8841 len: 115 ! Aa004881 Human polypeptide SEQ ID NO 18773.

24: ITPHP (R,K) {20,20} KKKKK
25: TPHPK (K) {20} KKKKK
26: PHPKK (K) {20} KKKKK
27: HPKKK (K) {20} KKKKG
28: PKKKK (K) {20} KKKGG
29: KKKKK (K) {20} KKGGA
30: KKKKK (K) {20} KKGAL
31: KKKKK (K) {20} GGALK

AA004917 ck: 3999 len: 103 ! Aa004917 Human polypeptide SEQ ID NO 18809.

1: (R,K) {20,20} KKKKKKKKKKKKKKKKKKK GGGPL

AA004928 ck: 23 len: 43 ! Aa004928 Human polypeptide SEQ ID NO 18820.

19: FKKEE (R,K) {20,20} KKKKKKKKKKKKKKKKKKK GGGGP (K) {20}

1

AA004969 ck: 4345 len: 57 ! Aa004969 Human polypeptide SEQ ID NO 18861.

(R,K){20,20}

(K){20}

26: LMGTG KKKKKKKKKKKKKKKKKKKKK

(K){20}

27: MGTSG KKKKKKKKKKKKKKKKKKKKK

(K){20}

28: GTSRK KKKKKKKKKKKKKKKKKKKKK

(K){20}

29: TSKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

30: SKRKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

31: KKKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

32: KKKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

33: KKKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

34: KKKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

35: KKKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

36: KKKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

37: KKKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

38: KKKKK KKKKKKKKKKKKKKKKKKKKK

AA005000 ck: 5847 len: 38 ! Aa005000 Human polypeptide SEQ ID NO 18892.

(R,K){20,20}

(K){20}

5: SGRS KKKKKKKKKKKKKKKKKKKKK

(K){20}

6: SGRSK KKKKKKKKKKKKKKKKKKKKK

(K){20}

7: GRSKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

8: RSKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

9: SKKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

10: KKKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

11: KKKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

12: KKKKK KKKKKKKKKKKKKKKKKKKKK

(K){20}

13: KKKKK KKKKKKKKKKKKKKKKKKKKK

1

14: KKKKK (K){20} KKKKKKKKKKKKKKKKKKK KRGGR

(K){20}

15: KKKKK KKKKKKKKKKKKKKKKKKKKK RGR

(R,K){20}

16: KKKKK KKKKKKKKKKKKKKKKKKKKK GGR

AA005081 ck: 2260 len: 40 ! Aa005081 Human polypeptide SEQ ID NO 18973.

(R,K){20,20}

(K){20}

9: SPASP KKKKKKKKKKKKKKKKKKKKK KKKK

(K){20}

10: RASPK KKKKKKKKKKKKKKKKKKKKK KKKK

(K){20}

11: ASPKK KKKKKKKKKKKKKKKKKKKKK KKKK

(K){20}

12: SPKKK KKKKKKKKKKKKKKKKKKKKK KKKK

(K){20}

13: PKKKK KKKKKKKKKKKKKKKKKKKKK KKKK

(K){20}

14: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKK

(K){20}

15: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKK

(K){20}

16: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKK

(K){20}

17: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKK

(K){20}

18: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKK

AA005130 ck: 6597 len: 116 ! Aa005130 Human polypeptide SEQ ID NO 19022.

(R,K){20,20}

(K){20}

14: ETPFQ KKKKKKKKKKKKKKKKKKKKK KRGGR

(K){20}

15: TPFQK KKKKKKKKKKKKKKKKKKKKK RGGGF

(R,K){20}

16: PFOKK KKKKKKKKKKKKKKKKKKKKK GGGFL

AA005191 ck: 4399 len: 85 ! Aa005191 Human polypeptide SEQ ID NO 19083.

(R,K){20,20}

(R,K){20}

30: GYIQL KKKKKKKKKKKKKKKKKKKKK GGAFF

AA005260 ck: 3618 len: 55 ! Aa005260 Human polypeptide SEQ ID NO 19152.

(R,K){20,20}

(K){20}

15: PHHQK KKKKKKKKKKKKKKKKKKKKK KKKK

(K){20}

16: HGHQK KKKKKKKKKKKKKKKKKKKKK KKKK

1 AAO05665 ck: 432 len: 28 ! Aao05665 Human polypeptide SEQ ID NO 19557.
(R,K){20,20}
(K){20}
6: DFLIQ KKKKKKKKKKKKKKKKKKKKK RGG
(R,K){20}
7: FLIQK KKKKKKKKKKKKKKKKKKKR GG

1 AAO06186 ck: 998 len: 88 ! Aao06186 Human polypeptide SEQ ID NO 20078.
(R,K){20,20}
(K){20}
28: SLIPK KKKKKKKKKKKKKKKKKKKKK KKKKG
(K){20}
29: LIPK KKKKKKKKKKKKKKKKKKKKK KKKGG
(K){20}
30: LIPK KKKKKKKKKKKKKKKKKKKKK KKKGG
(K){20}
31: PKKK KKKKKKKKKKKKKKKKKKKKK KGGGG
(K){20}
32: XKKK KKKKKKKKKKKKKKKKKKKKK GGGGF

1 AAO06357 ck: 4679 len: 52 ! Aao06357 Human polypeptide SEQ ID NO 20249.
(R,K){20,20}
(K){20}
9: FTCLI KKKKKKKKKKKKKKKKKKKKK KKKIK
(K){20}
10: TCLI KKKKKKKKKKKKKKKKKKKKK KKKIX
(K){20}
11: CLIK KKKKKKKKKKKKKKKKKKKKK KIKKK
(K){20}
12: LIKK KKKKKKKKKKKKKKKKKKKKK IKKKK

1 AAO06429 ck: 2585 len: 71 ! Aao06429 Human polypeptide SEQ ID NO 20321.
(R,K){20,20}
(K){20}
29: VIIMK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
30: IIMK KKKKKKKKKKKKKKKKKKKKK KKKKG
(K){20}
31: IMKK KKKKKKKKKKKKKKKKKKKKK KKKGR
(K){20}
32: MKKK KKKKKKKKKKKKKKKKKKKKK KKKRG
(K){20}
33: XKKK KKKKKKKKKKKKKKKKKKKKK KKKRG
(K){20}
34: KKKK KKKKKKKKKKKKKKKKKKKKK KRGGA

1 AAO06922 ck: 7296 len: 111 ! Aao06922 Human polypeptide SEQ ID NO 20814.
(R,K){20,20}
(K){20}
13: HSEL KKKKKKKKKKKKKKKKKKKKK GGGPK

1 AAO07241 ck: 3134 len: 40 ! Aao07241 Human polypeptide SEQ ID NO 21133.
(R,K){20,20}
(K){20}
4: YFP KKKKKKKKKKKKKKKKKKKKK RKKKX
(R,K){20}
5: YFPK KKKKKKKKKKKKKKKKKKKKK KKKKK
(R,K){20}
6: YFPK KKKKKKKKKKKKKKKKKKKKK KKKKG
(R,K){20}
7: FPKK KKKKKKKKKKKKKKKKKKKKK KKKGG
(R,K){20}
8: PKKK KKKKKKKKKKKKKKKKKKKKK XKKGG

1 AAO07354 ck: 6513 len: 133 ! Aao07354 Human polypeptide SEQ ID NO 21246.
(R,K){20,20}
(K){20}
36: KYHL KKKKKKKKKKKKKKKKKKKKK KKKGG
(K){20}
37: XYHL KKKKKKKKKKKKKKKKKKKKK KKKGP
(K){20}
38: YHLK KKKKKKKKKKKKKKKKKKKKK KGGPL
(K){20}
39: HLKK KKKKKKKKKKKKKKKKKKKKK GGPKL

1 AAO07410 ck: 693 len: 80 ! Aao07410 Human polypeptide SEQ ID NO 21302.
(R,K){20,20}
(K){20}
56: HNPI KKKKKKKKKKKKKKKKKKKKK RGGGF
(R,K){20}
57: KPIK KKKKKKKKKKKKKKKKKKKKK GGGF

1 AAO07412 ck: 4623 len: 166 ! Aao07412 Human polypeptide SEQ ID NO 21304.
(R,K){20,20}
(K){20}
93: VNTQ KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
94: NTQK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
95: TQKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
96: QKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
97: XKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
98: KKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
99: KKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
100: KKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
101: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
102: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
103: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
104: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
105: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
106: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
107: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

AA007505 ck: 3528 len: 72 ! Aa007505 Human polypeptide SEQ ID NO 21397.

1 (R, K) {20, 20}

14: HLVEA KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

15: LVEAK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

16: VEAkk KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

17: EAAKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

18: AKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

19: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

20: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

21: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

22: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

23: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

AA007509 ck: 4998 len: 140 ! Aa007509 Human polypeptide SEQ ID NO 21401.

1 (R, K) {20, 20}

17: KVVKE KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

18: VKKEK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

19: KKEKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

20: XEKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

21: EKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

22: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

23: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

24: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

25: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

26: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

27: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

28: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

29: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

30: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

31: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

32: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

AA007594 ck: 7271 len: 93 ! Aa007594 Human polypeptide SEQ ID NO 21486.

1 (R, K) {20, 20}

9: FLLLG KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

10: LLLGK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

11: LLGKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

12: LGKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

13: GKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

AA007607 ck: 2434 len: 72 ! Aa007607 Human polypeptide SEQ ID NO 21499.

1 (R, K) {20, 20}

21: KKKKE KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

48: GLFXE KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

AA007610 ck: 5270 len: 74 ! Aa007610 Human polypeptide SEQ ID NO 21502.

1 (R, K) {20, 20}

19: TELTI KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

20: ELTIK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

21: LTIKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKK
22: TTIKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKK
23: IKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKGG
24: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKGG
25: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKGGA
26: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} GGGA

AAO07755 ck: 6069 len: 90 ! Aao07755 Human polypeptide SEQ ID NO 21647.

35: KNOSS KKKKKKKKKKKKKKKKKKK (R,K) {20,20} KKKK
36: NQSSK KKKKKKKKKKKKKKKKKKK (K) {20} KKKK
37: QSSKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKK
38: SSKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKK
39: SKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKGG
40: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKGP
41: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKGPL
42: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} GGPL

AAO07762 ck: 5115 len: 41 ! Aao07762 Human polypeptide SEQ ID NO 21654.

18: IPSLK KKKKKKKKKKKKKKKKKKK (R,K) {20,20} KKKK
19: PSLKK KKKKKKKKKKKKKKKKKKK (K) {20} KKK
20: SLKKK KKKKKKKKKKKKKKKKKKK (K) {20} KK
21: LKKKK KKKKKKKKKKKKKKKKKKK (K) {20} K
22: XKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKK

AAO07775 ck: 9827 len: 90 ! Aao07775 Human polypeptide SEQ ID NO 21667.

46: PHKPE KKKKKKKKKKKKKKKKKKK (R,K) {20,20} KKKK
(K) {20}

47: HXPEK KKKKKKKKKKKKKKKKKKK (K) {20} KKKGG
48: XPEKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKGG
49: PEKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKGGA
50: EKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKGAP
51: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} GGAPK

AAO07806 ck: 1959 len: 99 ! Aao07806 Human polypeptide SEQ ID NO 21698.

41: KLHCI KKKKKKKKKKKKKKKKKKK (R,K) {20,20} KKKK
42: LHCIC KKKKKKKKKKKKKKKKKKK (K) {20} KKKK
43: HCICK KKKKKKKKKKKKKKKKKKK (K) {20} KKKK
44: CIKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKL
45: IKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKLW
46: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KKLWG
47: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} KLWGG
48: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20} LMGGG

AAO07872 ck: 8003 len: 135 ! Aao07872 Human polypeptide SEQ ID NO 21764.

52: TRKKQ KKKKKKKKKKKKKKKKKKK (R,K) {20,20} GGGLL
AAO07874 ck: 8007 len: 111 ! Aao07874 Human polypeptide SEQ ID NO 21766.
90: EEPIS KKKKKKKKKKKKKKKKKKK (R,K) {20,20} KG
91: EPSIK KKKKKKKKKKKKKKKKKKK (K) {20} G

AAO07876 ck: 8130 len: 66 ! Aao07876 Human polypeptide SEQ ID NO 21768.

17: IYSFE KKKKKKKKKKKKKKKKKKK (R,K) {20,20} KKKK
18: YSFEK KKKKKKKKKKKKKKKKKKK (K) {20} KKKK
19: SFEKK KKKKKKKKKKKKKKKKKKK (K) {20} KKKKQ

20: PEEKK (K) {20}
21: EKKKK (K) {20}
22: KKKKK (K) {20}
23: KKKKK (K) {20}

AA007976 ck: 6471 len: 97 i Aa007976 Human polypeptide SEQ ID NO 21868.

43: PPIST (R,K) {20,20}
44: PISTK (K) {20}
45: ISTKK (K) {20}
46: STKKK (K) {20}
47: TKKKK (K) {20}
48: KKKKK (K) {20}
49: KKKKK (K) {20}
50: KKKKK (K) {20}
51: KKKKK (K) {20}
52: KKKKK (K) {20}
53: KKKKK (K) {20}
54: KKKKK (K) {20}
55: KKKKK (K) {20}

AA007981 ck: 9551 len: 81 i Aa007981 Human polypeptide SEQ ID NO 21873.

54: KPCIY (R,K) {20,20}
55: PCIYK (K) {20}
56: CIYKK (K) {20}
57: IYKKK (K) {20}
58: YKKKK (K) {20}

59: KKKKK (K) {20}
60: KKKKK (K) {20}
61: KKKKK (K) {20}
62: KKKKK (K) {20}

AA008018 ck: 9768 len: 48 i Aa008018 Human polypeptide SEQ ID NO 21910.

24: KISQV (R,K) {20,20}
25: ISQVR (K) {20}
26: SQVRK (K) {20}
27: QVRKK (K) {20}
28: VRKKK (K) {20}
29: RKKKK (K) {20}

AA008121 ck: 5381 len: 86 i Aa008121 Human polypeptide SEQ ID NO 22013.

20: HFGLL (R,K) {20,20}

20: HFGLL (K) {20}

AA008232 ck: 782 len: 104 i Aa008232 Human polypeptide SEQ ID NO 22124.

49: ISQON (R,K) {20,20}
50: SQONK (K) {20}

AA008450 ck: 6228 len: 31 i Aa008450 Human polypeptide SEQ ID NO 22342.

5: NNSA (R,K) {20,20}
6: NNSAK (K) {20}
7: NSAKK (K) {20}

AA008458 ck: 1813 len: 59 i Aa008458 Human polypeptide SEQ ID NO 22350.

8: LGWBE (R,K) {20,20}

9: GWEK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
10: GWEK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
11: WEKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
12: EKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
13: KKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

AA008460 ck: 5334 len: 62 ! Aa008460 Human polypeptide SEQ ID NO 22352.

(R,K) {20,20}

22: KEFST KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

23: EFSTK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

24: FSTKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

25: STYKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

26: TKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

27: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

28: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

29: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

30: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

31: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

32: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

33: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

34: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

AA008466 ck: 8728 len: 107 ! Aa008466 Human polypeptide SEQ ID NO 22358.

(R,K) {20,20}

57: FMGGV KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

58: WGGVY KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

59: GGVKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

60: GVKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

61: VKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

62: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

63: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(R,K) {20}

64: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(R,K) {20}

65: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(R,K) {20}

AA008580 ck: 5164 len: 62 ! Aa008580 Human polypeptide SEQ ID NO 22472.

(R,K) {20,20}

30: PILLK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

AA008591 ck: 900 len: 46 ! Aa008591 Human polypeptide SEQ ID NO 22483.

(R,K) {20,20}

14: LLSQK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

AA008609 ck: 5151 len: 74 ! Aa008609 Human polypeptide SEQ ID NO 22501.

(R,K) {20,20}

18: KCVIL KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

19: CVILK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

AA008612 ck: 557 len: 46 ! Aa008612 Human polypeptide SEQ ID NO 22504.

(R,K) {20,20}

20: TFCIM KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

21: FCIMK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

22: CIMKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

23: IMKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

AA008616 ck: 4555 len: 41 ! Aa008616 Human polypeptide SEQ ID NO 22508.

(R,K) {20,20}

6: HCALP KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

7: CALPK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

8: ALPKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

9: LPKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

10: PKKKK KKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

11: KKKK (K) {20} KKKK
12: KKKK (K) {20} KKKK
13: KKKK (K) {20} KKKK
14: KKKK (K) {20} KKKK
15: KKKK (K) {20} KKKK
16: KKKK (K) {20} KKKK
17: KKKK (K) {20} KKKK
18: KKKK (K) {20} KKKK
19: KKKK (K) {20} KKKK
20: KKKK (K) {20} KKKK
21: KKKK (K) {20} K
22: KKKK (K) {20} KKKK

AA008623 ck: 9243 len: 119 ! Aa008623 Human polypeptide SEQ ID NO 22515.
(R,K) {20,20}
(R,K) {20}

21: DSKQE KKKKKKKKKKKKKKKR GGAFK

11: EGNMW KKKKKKKKKKKKKKKK KKKK
12: GNNWK KKKKKKKKKKKKKKKK KKKK
13: NNWK KKKKKKKKKKKKKKKK KKKK
14: NNWK KKKKKKKKKKKKKKKK KKKK
15: WKKK KKKKKKKKKKKKKKKK KKKK
16: KKKK KKKKKKKKKKKKKKKK KKKK

AA008631 ck: 6430 len: 135 ! Aa008631 Human polypeptide SEQ ID NO 22523.
(R,K) {20,20}
(K) {20}

30: TPSRA KKKKKKKKKKKKKKKK KKKK

31: PSRAK KKKKKKKKKKKKKKKK KKKK
32: SPRAK KKKKKKKKKKKKKKKK KKKK
33: PAKK KKKKKKKKKKKKKKKK KKKK
34: AKKK KKKKKKKKKKKKKKKK KKKK
35: KKKK KKKKKKKKKKKKKKKK KKKK
36: KKKK KKKKKKKKKKKKKKKK KKKK
37: KKKK KKKKKKKKKKKKKKKK KKKK
38: KKKK KKKKKKKKKKKKKKKK KKKK
39: KKKK KKKKKKKKKKKKKKKK KKKK
40: KKKK KKKKKKKKKKKKKKKK KKKK
41: KKKK KKKKKKKKKKKKKKKK KKKK
42: KKKK KKKKKKKKKKKKKKKK KKKK
43: KKKK KKKKKKKKKKKKKKKK KKKK
44: KKKK KKKKKKKKKKKKKKKK KKKK

AA008644 ck: 3202 len: 80 ! Aa008644 Human polypeptide SEQ ID NO 22536.
(R,K) {20,20}
(K) {20}

2: E KKKKKKKKKKKKKKKK KKKK
3: EK KKKKKKKKKKKKKKKK KKKK
4: EKK KKKKKKKKKKKKKKKK KKKK
5: EKK KKKKKKKKKKKKKKKK KKKK
6: EKKK KKKKKKKKKKKKKKKK KKKK
7: KKKK KKKKKKKKKKKKKKKK KKKK
8: KKKK KKKKKKKKKKKKKKKK KKKK
9: KKKK KKKKKKKKKKKKKKKK KKKK
10: KKKK KKKKKKKKKKKKKKKK KKKK

11: KKKK (K) {20} KKKK
12: KKKK (K) {20} KKKK
13: KKKK (K) {20} KKKK
14: KKKK (K) {20} KKKK
15: KKKK (K) {20} KKKK
16: KKKK (K) {20} KKKK
17: KKKK (K) {20} KKKK
18: KKKK (K) {20} KKKK
19: KKKK (K) {20} KKKK

AA008653 ck: 6448 len: 63 1 Aa008653 Human polypeptide SEQ ID NO 22545.

1

12: PIKLT (R,K) {20,20} KKKK
13: IKLT (K) {20} KKKK
14: KLT (K) {20} KKKK
15: LTKK (K) {20} KKKK
16: TKKK (K) {20} KKKK
17: KKKK (K) {20} KKKK
18: KKKK (K) {20} KKKK
19: KKKK (K) {20} KKKK
20: KKKK (R,K) {20} KKKK

AA008686 ck: 6936 len: 81 1 Aa008686 Human polypeptide SEQ ID NO 22578.

1

23: HPVNH (R,K) {20,20} KKKK
24: PVNH (K) {20} KKKK
25: VNHK (K) {20} KKKK
26: NHKK (K) {20} KKKK

27: HKKK (K) {20} KKKK
28: KKKK (K) {20} KKKK
29: KKKK (K) {20} KKKK
30: KKKK (K) {20} KKKK
31: KKKK (K) {20} KKKK
32: KKKK (K) {20} KKKK
33: KKKK (K) {20} KKKK
34: KKKK (K) {20} KKKK
35: KKKK (K) {20} KKKK
36: KKKK (K) {20} KKKK
37: KKKK (K) {20} KKKK

AA008701 ck: 588 len: 46 1 Aa008701 Human polypeptide SEQ ID NO 22593.

1

7: INANS (R,K) {20,20} KKKK
8: NANSK (K) {20} KKKK
9: ANSK (K) {20} KKKK
10: NSKK (K) {20} KKKK
11: SKKK (K) {20} KKKK
12: KKKK (K) {20} KKKK
13: KKKK (K) {20} KKKK

AA008707 ck: 9359 len: 48 1 Aa008707 Human polypeptide SEQ ID NO 22599.

1

13: TAVTQ (R,K) {20,20} KKKK
14: AVTQK (K) {20} KKKK
15: VTQKK (K) {20} KKKK

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1 AA008708 ck: 9654 len: 32 ! Aa008708 Human polypeptide SEQ ID NO 22600.
      (R,K){20,20}
      (K){20}
8: SMTEP KKKKKKKKKKKKKKKKKKK KKKKK
9: MTEPK KKKKKKKKKKKKKKKKKKK KKKK
      (K){20}
10: TPEPK KKKKKKKKKKKKKKKKKKK KKK
      (K){20}
11: FPKKK KKKKKKKKKKKKKKKKKKK KK
      (K){20}
12: PKKKK KKKKKKKKKKKKKKKKKKK K
      (K){20}
13: KKKKK KKKKKKKKKKKKKKKKKKK

1 AA008717 ck: 4688 len: 99 ! Aa008717 Human polypeptide SEQ ID NO 22609.
      (R,K){20,20}
      (K){20}
80: SPFLI KKKKKKKKKKKKKKKKKKK

1 AA008772 ck: 1431 len: 105 ! Aa008772 Human polypeptide SEQ ID NO 22664.
      (R,K){20,20}
      (K){20}
23: IMNYL KKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
24: MNYLK KKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
25: NYLKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
26: YLKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
27: LKKKK KKKKKKKKKKKKKKKKKKK KKKGG
      (K){20}
28: KKKKK KKKKKKKKKKKKKKKKKKK KKGCG
      (K){20}
29: KKKKK KKKKKKKKKKKKKKKKKKK KGGGP
      (K){20}
30: KKKKK KKKKKKKKKKKKKKKKKKK GGGPF

AA008774 ck: 2529 len: 71 ! Aa008774 Human polypeptide SEQ ID NO 22666.
      (R,K){20,20}
      (K){20}
16: RTIKL KKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
17: TIKLK KKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
18: IKLKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
19: KLKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
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20: LKKKK KKKKKKKKKKKKKKKKKKK KKKGG
      (K){20}
21: KKKKK KKKKKKKKKKKKKKKKKKK KKGGP
      (K){20}
22: KKKKK KKKKKKKKKKKKKKKKKKK KGGPP
      (K){20}
23: KKKKK KKKKKKKKKKKKKKKKKKK GGGPF

1 AA008817 ck: 529 len: 28 ! Aa008817 Human polypeptide SEQ ID NO 22709.
      (R,K){20,20}
      (K){20}
3: AX KKKKKKKKKKKKKKKKKKK NIIMG

1 AA008820 ck: 1888 len: 84 ! Aa008820 Human polypeptide SEQ ID NO 22712.
      (R,K){20,20}
      (K){20}
28: CMTFS KKKKKKKKKKKKKKKKKKK RGGGF
      (R,K){20}
29: MTFPS KKKKKKKKKKKKKKKKKKK GGGFI

1 AA008841 ck: 3529 len: 74 ! Aa008841 Human polypeptide SEQ ID NO 22733.
      (R,K){20,20}
      (K){20}
3: PQ KKKKKKKKKKKKKKKKKKK KMGCG
      (K){20}
4: PQK KKKKKKKKKKKKKKKKKKK WGGGF

1 AA008857 ck: 2816 len: 49 ! Aa008857 Human polypeptide SEQ ID NO 22749.
      (R,K){20,20}
      (K){20}
25: PPPTS KKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
26: PPTSK KKKKKKKKKKKKKKKKKKK KKKK
      (K){20}
27: PTSKK KKKKKKKKKKKKKKKKKKK KKK
      (K){20}
28: TSKKK KKKKKKKKKKKKKKKKKKK KK
      (K){20}
29: SKKKK KKKKKKKKKKKKKKKKKKK K
      (K){20}
30: KKKKK KKKKKKKKKKKKKKKKKKK

1 AA008913 ck: 3132 len: 52 ! Aa008913 Human polypeptide SEQ ID NO 22805.
      (R,K){20,20}
      (K){20}
9: IIKSF KKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
10: IKSFK KKKKKKKKKKKKKKKKKKK KKKKK
      (K){20}
11: KSFKK KKKKKKKKKKKKKKKKKKK KKKKK
```

12: SPKKK (K) {20}
13: FKKKK (K) {20}
14: KKKKK (K) {20}
15: KKKKK (K) {20}
16: KKKKK (K) {20}
17: KKKKK (K) {20}
18: KKKKK (K) {20}
19: KKKKK (K) {20}
20: KKKKK (K) {20}
21: KKKKK (K) {20}
22: KKKKK (K) {20}
23: KKKKK (K) {20}
24: KKKKK (K) {20}
25: KKKKK (K) {20}
26: KKKKK (K) {20}
27: KKKKK (K) {20}
28: KKKKK (K) {20}
29: KKKKK (K) {20}
30: KKKKK (K) {20}
31: KKKKK (K) {20}

AAO08914 ck: 4641 len: 25 ! Aao08914 Human polypeptide SEQ ID NO 22806.
(R,K) {20,20}
(R,K) {20}

6: ILMPX KKKKKKKKKKKKKKKKKKKR

AAO08943 ck: 5770 len: 75 ! Aao08943 Human polypeptide SEQ ID NO 22835.
(R,K) {20,20}
(R,K) {20}

8: SEWAA KKKKKKKKKKKKKKKKKKK KKKIF

9: EWAAR (K) {20}
10: WPAAR (K) {20}
11: AAKKK (K) {20}

AAO08994 ck: 2695 len: 29 ! Aao08994 Human polypeptide SEQ ID NO 22886.
(R,K) {20,20}
(K) {20}

9: NLPSS KKKKKKKKKKKKKKKKKKK K
10: LPSSK KKKKKKKKKKKKKKKKKKK

AAO08995 ck: 6270 len: 26 ! Aao08995 Human polypeptide SEQ ID NO 22887.
(R,K) {20,20}
(K) {20}

6: MAAPP KKKKKKKKKKKKKKKKKKK I

AAO09001 ck: 6400 len: 26 ! Aao09001 Human polypeptide SEQ ID NO 22893.
(R,K) {20,20}
(K) {20}

5: LTSS KKKKKKKKKKKKKKKKKKK KK
6: LTSSK KKKKKKKKKKKKKKKKKKK K
7: TSSKK KKKKKKKKKKKKKKKKKKK

AAO09016 ck: 4378 len: 119 ! Aao09016 Human polypeptide SEQ ID NO 22908.
(R,K) {20,20}
(R,K) {20}

10: EEEER RKKKKRRRRRRKKKKKK ILRQK

AAO09057 ck: 3712 len: 58 ! Aao09057 Human polypeptide SEQ ID NO 22949.
(R,K) {20,20}
(K) {20}

5: MILN KKKKKKKKKKKKKKKKKKK KKKKK
6: MILNK KKKKKKKKKKKKKKKKKKK KKKGG
7: ILNKK KKKKKKKKKKKKKKKKKKK KKKGG
8: LNKKK KKKKKKKKKKKKKKKKKKK KKKGG
9: NKKKK KKKKKKKKKKKKKKKKKKK KGGGP
10: KKKKK KKKKKKKKKKKKKKKKKKK GGGPF

AAO09066 ck: 2645 len: 29 ! Aao09066 Human polypeptide SEQ ID NO 22958.
(R,K) {20,20}

4: AQQ ^{(K) {20}} KKKKKKKKKKKKKKKKKKK KKKKK
5: AQQK ^{(K) {20}} KKKKKKKKKKKKKKKKKKK KKKKK
6: AQQKK ^{(K) {20}} KKKKKKKKKKKKKKKKKKK KKKKK
7: QQQKK ^{(K) {20}} KKKKKKKKKKKKKKKKKKK KKK
8: QKKKK ^{(K) {20}} KKKKKKKKKKKKKKKKKKK KK
9: KKKKK ^{(K) {20}} KKKKKKKKKKKKKKKKKKK K
10: KKKKK ^{(K) {20}} KKKKKKKKKKKKKKKKKKK

AA009072 ck: 8432 len: 42 ! Aa009072 Human polypeptide SEQ ID NO 22364.
^{(R,K) {20,20}}
^{(K) {20}}
11: LIFTL KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
12: IFTLK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
13: FTLLK KKKKKKKKKKKKKKKKKKK KKKKK
14: TLKKK ^{(K) {20}} KKKKKKKKKKKKKKKKKKK KKKKK

AA009077 ck: 9524 len: 87 ! Aa009077 Human polypeptide SEQ ID NO 22369.
^{(R,K) {20,20}}
^{(K) {20}}
26: RPLLT KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
27: FLILT KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
28: LLTKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
29: LTKKK KKKKKKKKKKKKKKKKKKK KKKKK
30: TKKKK ^{(K) {20}} KKKKKKKKKKKKKKKKKKK KKKKK

AA009162 ck: 3841 len: 100 ! Aa009162 Human polypeptide SEQ ID NO 23054.
^{(R,K) {20,20}}
^{(K) {20}}
11: KTLFQ KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
12: TLFQK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
13: LFQKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
14: FQKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}

15: QKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
16: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
17: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
18: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
19: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
20: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
21: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
22: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
23: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
24: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
25: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
26: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
27: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
28: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
29: KKKKK ^{(R,K) {20}} KKKKKKKKKKKKKKKKKKK KKKKK

AA009258 ck: 7988 len: 42 ! Aa009258 Human polypeptide SEQ ID NO 23150.
^{(R,K) {20,20}}
^{(K) {20}}
7: LFFCP KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
8: LFPCP KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
9: FPCPK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
10: CPCKP KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
11: PKKKK KKKKKKKKKKKKKKKKKKK KKKKK
^{(K) {20}}
12: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

AA009269 ck: 7553 len: 106 ! Aa009269 Human polypeptide SEQ ID NO 23161.
^{(R,K) {20,20}}
^{(R,K) {20}}
39: NTGML KKKKKKKKKKKKKKKKKKKR GGPFLL

1 AAO09457 ck: 9658 len: 32 ! Aao09457 Human polypeptide SEQ ID NO 23349.
(R,K){20,20}
(K){20}
9: ALVPQ KKKKKKKKKKKKKKKKKKKKK NIKI

1 AAO09819 ck: 4709 len: 80 ! Aao09819 Human polypeptide SEQ ID NO 23711.
(R,K){20,20}
(K){20}
54: CPEVX KKKKKKKKKKKKKKKKKKKKK EKGSG

1 AAO10447 ck: 6342 len: 50 ! Aao10447 Human polypeptide SEQ ID NO 24339.
(R,K){20,20}
(K){20}
25: RPLKL KKKKKKKKKKKKKKKKKKKKK RKKKK
(R,K){20}
26: PLKLK KKKKKKKKKKKKKKKKKKKKK KKKKI
(R,K){20}
27: LKLKK KKKKKKKKKKKKKKKKKKKKK KKKI
(R,K){20}
28: KLKKK KKKKKKKKKKKKKKKKKKKKK KKI
(R,K){20}
29: LKKKK KKKKKKKKKKKKKKKKKKKKK KI
(R,K){20}
30: KKKKK KKKKKKKKKKKKKKKKKKKKK I

1 AAO10451 ck: 3955 len: 82 ! Aao10451 Human polypeptide SEQ ID NO 24343.
(R,K){20,20}
(K){20}
59: SRASP KKKKKKKKKKKKKKKKKKKKK ARGG

1 AAO10467 ck: 7542 len: 116 ! Aao10467 Human polypeptide SEQ ID NO 24359.
(R,K){20,20}
(K){20}
57: KCEFM KKKKKKKKKKKKKKKKKKKKK KIGGG
(K){20}
58: CEFMK KKKKKKKKKKKKKKKKKKKKK IGSGA

1 AAO10564 ck: 9156 len: 98 ! Aao10564 Human polypeptide SEQ ID NO 24456.
(R,K){20,20}
(K){20}
18: NLTLT KKKKKKKKKKKKKKKKKKKKK KKEMP
(K){20}
19: LTLTK KKKKKKKKKKKKKKKKKKKKK KEMPV
(K){20}
20: LTLKK KKKKKKKKKKKKKKKKKKKKK EMPVK

1 AAO10608 ck: 9106 len: 67 ! Aao10608 Human polypeptide SEQ ID NO 24500.
(R,K){20,20}
(K){20}
42: CRUSE KKKKKKKKKKKKKKKKKKKKK KKKKD

43: RLSEK KKKKKKKKKKKKKKKKKKKKK KKDS
(K){20}
44: LSEKK KKKKKKKKKKKKKKKKKKKKK KKDS
(K){20}
45: SEKKK KKKKKKKKKKKKKKKKKKKKK KDS
(K){20}
46: EKKKK KKKKKKKKKKKKKKKKKKKKK DS

1 AAO10638 ck: 621 len: 56 ! Aao10638 Human polypeptide SEQ ID NO 24530.
(R,K){20,20}
(K){20}
33: CEPQP KKKKKKKKKKKKKKKKKKKKK KARG
(K){20}
34: EPQPK KKKKKKKKKKKKKKKKKKKKK ARG

1 AAO10786 ck: 7349 len: 65 ! Aao10786 Human polypeptide SEQ ID NO 24678.
(R,K){20,20}
(K){20}
24: CPXFS KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
25: PYFSK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
26: XFSKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
27: FSKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
28: SKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
29: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
30: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
31: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKV
(K){20}
32: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKVF
(K){20}
33: KKKKK KKKKKKKKKKKKKKKKKKKKK KKVFF
(K){20}
34: KKKKK KKKKKKKKKKKKKKKKKKKKK KVFFF
(K){20}
35: KKKKK KKKKKKKKKKKKKKKKKKKKK VFFFE

1 AAO10852 ck: 5147 len: 41 ! Aao10852 Human polypeptide SEQ ID NO 24744.
(R,K){20,20}
(K){20}
20: FSRXM KKKKKKKKKKKKKKKKKKKKK KK
(K){20}
21: SRXMK KKKKKKKKKKKKKKKKKKKKK K
(K){20}

22: RMXKK KKKKKKKKKKKKKKKKKKK

AAO10853 ck: 444 len: 98 ! Aao10853 Human polypeptide SEQ ID NO 24745.

(R,K) {20,20}

(K) {20}

14: LRAHL KKKKKKKKKKKKKKKKKKK

(K) {20}

15: RAHLK KKKKKKKKKKKKKKKKKKK

(K) {20}

16: AHLK KKKKKKKKKKKKKKKKKKK

(K) {20}

17: HLKK KKKKKKKKKKKKKKKKKKK

(K) {20}

18: LKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

19: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

20: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

21: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

22: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

23: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

24: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

25: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

26: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

27: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

28: KKKK KKKKKKKKKKKKKKKKKKK

(R,K) {20}

29: KKKK KKKKKKKKKKKKKKKKKKK

AAO10859 ck: 2690 len: 70 ! Aao10859 Human polypeptide SEQ ID NO 24751.

(R,K) {20,20}

(K) {20}

44: XGILK KKKKKKKKKKKKKKKKKKK

(K) {20}

45: GITEK KKKKKKKKKKKKKKKKKKK

AAO10933 ck: 849 len: 69 ! Aao10933 Human polypeptide SEQ ID NO 24825.

(R,K) {20,20}

(K) {20}

33: FINTE KKKKKKKKKKKKKKKKKKK

(K) {20}

34: INTEK KKKKKKKKKKKKKKKKKKK

(K) {20}

35: NTEKK KKKKKKKKKKKKKKKKKKK

(K) {20}

36: TEKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

37: EKKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

38: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

39: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

40: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

41: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

42: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

43: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

44: KKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

45: KKKK KKKKKKKKKKKKKKKKKKK

AAO10973 ck: 824 len: 51 ! Aao10973 Human polypeptide SEQ ID NO 24865.

(R,K) {20,20}

(R,K) {20}

16: SYFXM KKKKKKKKKKKKKKKKKKK

(K) {20}

17: YFXMR KKKKKKKKKKKKKKKKKKK

AAO10983 ck: 694 len: 51 ! Aao10983 Human polypeptide SEQ ID NO 24875.

(R,K) {20,20}

(K) {20}

24: HPILX KKKKKKKKKKKKKKKKKKK

(K) {20}

25: PILXK KKKKKKKKKKKKKKKKKKK

(K) {20}

26: ILXKK KKKKKKKKKKKKKKKKKKK

(K) {20}

27: LXXKK KKKKKKKKKKKKKKKKKKK

(K) {20}

28: XKKKK KKKKKKKKKKKKKKKKKKK

(K) {20}

29: KKKK KKKKKKKKKKKKKKKKKKK

AAO10995 ck: 2382 len: 122 ! Aao10995 Human polypeptide SEQ ID NO 24887.

(R,K) {20,20}

(K) {20}

22: PCGGS KKKKKKKKKKKKKKKKKKK

23: CKGSK KKKKKKKKKKKKKKKKKKK (K) {20}
24: KGSKK KKKKKKKKKKKKKKKKKKK (K) {20}
25: GSKKK KKKKKKKKKKKKKKKKKKK (K) {20}
26: SKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
27: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
28: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
AA010997 ck: 5315 len: 58 ! Aa010997 Human polypeptide SEQ ID NO 24889.
(R, K) {20, 20}
29: IXIIQ KKKKKKKKKKKKKKKKKKK (K) {20}
30: XIIOK KKKKKKKKKKKKKKKKKKK (K) {20}
31: IIOKK KKKKKKKKKKKKKKKKKKK (K) {20}
32: IOKKK KKKKKKKKKKKKKKKKKKK (K) {20}
33: OKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
34: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
35: KKKKK KKKKKKKKKKKKKKKKKKK (R, K) {20}
AA011002 ck: 8638 len: 82 ! Aa011002 Human polypeptide SEQ ID NO 24894.
(R, K) {20, 20}
5: FKRS KKKKKKKKKKKKKKKKKKK (R, K) {20}
38: XFKRS KKKKKKKKKKKKKKKKKKK (R, K) {20}
AA011033 ck: 3780 len: 49 ! Aa011033 Human polypeptide SEQ ID NO 24925.
(R, K) {20, 20}
10: OKKFI KKKKKKKKKKKKKKKKKKK (R, K) {20}
11: XKFIR KKKKKKKKKKKKKKKKKKK (K) {20}
AA011048 ck: 7868 len: 85 ! Aa011048 Human polypeptide SEQ ID NO 24940.
(R, K) {20, 20}
53: IISDP KKKKKKKKKKKKKKKKKKK (K) {20}
54: ISDPK KKKKKKKKKKKKKKKKKKK (K) {20}

55: SDPKK KKKKKKKKKKKKKKKKKKK (K) {20}
56: DPKKK KKKKKKKKKKKKKKKKKKK (K) {20}
57: PKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
58: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
59: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
60: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
61: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
62: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
63: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
64: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
65: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
66: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
AA011078 ck: 4186 len: 100 ! Aa011078 Human polypeptide SEQ ID NO 24970.
(R, K) {20, 20}
39: PPKPD KKKKKKKKKKKKKKKKKKK (K) {20}
40: PKPDK KKKKKKKKKKKKKKKKKKK (K) {20}
41: KPDKK KKKKKKKKKKKKKKKKKKK (K) {20}
42: PDKKK KKKKKKKKKKKKKKKKKKK (K) {20}
43: DKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
44: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
45: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
46: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
47: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
48: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}
49: KKKKK KKKKKKKKKKKKKKKKKKK (K) {20}

1

50: KKKK (K) {20} KKKKKKKKKKKKKKKKK KRGKG
51: KKKK (K) {20} KKKKKKKKKKKKKKKKK RGGGG
52: KKKK (R,K) {20} KKKKKKKKKKKKKKKR GGGGF
AA01124 ck: 5121 len: 61 ! Aa01124 Human polypeptide SEQ ID NO 25016.
(R,K) {20,20}
16: KYSFL KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
17: YSFLK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
18: SFLKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
19: FLKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
20: LKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
21: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
22: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
23: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
24: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
25: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
26: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
27: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
28: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
29: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
30: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
31: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
32: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
33: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
34: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
35: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

1

36: KKKK (K) {20} KKKKKKKKKKKKKKKKK KKKKK
(K) {20}
37: KKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
38: KKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
39: KKKK KKKKKKKKKKKKKKKKKKK KKK
(K) {20}
40: KKKK KKKKKKKKKKKKKKKKKKK KK
(K) {20}
41: KKKK KKKKKKKKKKKKKKKKKKK K
(K) {20}
42: KKKK KKKKKKKKKKKKKKKKKKK

AA01139 ck: 8807 len: 68 ! Aa01139 Human polypeptide SEQ ID NO 25031.

(R,K) {20,20}

(K) {20}

22: VCRFP KKKKKKKKKKKKKKKKKKK KMEGG

23: CRFPK KKKKKKKKKKKKKKKKKKK MEGGG

(K) {20}

1

AA01165 ck: 7203 len: 62 ! Aa01165 Human polypeptide SEQ ID NO 25057.

(R,K) {20,20}

(K) {20}

18: DASMV KKKKKKKKKKKKKKKKKKK KKKKK

19: ASMPK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

20: SMVKK KKKKKKKKKKKKKKKKKKK KKKKK

21: MPVKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

22: VKKKK KKKKKKKKKKKKKKKKKKK KKKKA

23: KKKKK KKKKKKKKKKKKKKKKKKK KKKAS

(K) {20}

24: KKKKK KKKKKKKKKKKKKKKKKKK KKAAS

25: KKKKK KKKKKKKKKKKKKKKKKKK KAASS

(K) {20}

26: KKKKK KKKKKKKKKKKKKKKKKKK ASSSQ

1

AA01209 ck: 6562 len: 96 ! Aa01209 Human polypeptide SEQ ID NO 25101.

(R,K) {20,20}

(K) {20}

63: LLAYK KKKKKKKKKKKKKKKKKKK KKPXY

64: LLAYK KKKKKKKKKKKKKKKKKKK KPYXL

(K) {20}

(K) {20}

65: LAYKK KKKKKKKKKKKKKKKKKKKKK PKYLS

AA011210 ck: 863 len: 70 1 Aa011210 Human polypeptide SEQ ID NO 25102.

(R,K){20,20}

(K){20}

31: DLCLC KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

32: LCLCK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

33: CLCKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

34: LCCKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

35: CKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

36: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

37: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

38: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

39: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

40: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

41: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

42: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

43: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

44: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

45: KKKKK KKKKKKKKKKKKKKKKKKKKK PGGGG

AA011214 ck: 9584 len: 68 1 Aa011214 Human polypeptide SEQ ID NO 25106.

(R,K){20,20}

(K){20}

19: NRVCX KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

20: RVCXK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

21: VCXKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

22: CXKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

23: XXXKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

24: KKKKK KKKKKKKKKKKKKKKKKKKKK RGGPL

25: KKKKK (R,K){20} KKKKKKKKKKKKKKKKKKKR GGPLK

AA011241 ck: 1100 len: 36 1 Aa011241 Human polypeptide SEQ ID NO 25133.

(R,K){20,20}

(K){20}

8: XOTLT KKKKKKKKKKKKKKKKKKKKK KAYTS

(K){20}

9: OTLTK KKKKKKKKKKKKKKKKKKKKK AYTSK

AA011248 ck: 5013 len: 88 1 Aa011248 Human polypeptide SEQ ID NO 25140.

(R,K){20,20}

(K){20}

51: TCLYX KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

52: CLYXK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

53: LYXKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

54: YXKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

55: XKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

56: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K){20}

57: KKKKK KKKKKKKKKKKKKKKKKKKKK KKGGA

(K){20}

58: KKKKK KKKKKKKKKKKKKKKKKKKKK KKGAF

(K){20}

59: KKKKK KKKKKKKKKKKKKKKKKKKKK GGAFK

AA011293 ck: 6490 len: 73 1 Aa011293 Human polypeptide SEQ ID NO 25185.

(R,K){20,20}

(K){20}

46: KXDYF KKKKKKKKKKKKKKKKKKKKK KKKGG

(K){20}

47: XDYFK KKKKKKKKKKKKKKKKKKKKK KKSOG

(K){20}

48: DYFKK KKKKKKKKKKKKKKKKKKKKK KSGGG

(K){20}

49: YFKKK KKKKKKKKKKKKKKKKKKKKK SGGGA

AA011324 ck: 6024 len: 93 1 Aa011324 Human polypeptide SEQ ID NO 25216.

(R,K){20,20}

(R,K){20}

37: ISLSA KKKKKKKKKKKKKKKKKKKKK KKKGG

(K){20}

38: SLSAR KKKKKKKKKKKKKKKKKKKKK KKGGG

(K){20}

39: LSARK KKKKKKKKKKKKKKKKKKKKK KGGGG

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40: SARRK KKKKKKKKKKKKKKKKKKKKKK GGGGV
      (K) {20}

1  AA011327 ck: 3052 len: 102 ! Aa011327 Human polypeptide SEQ ID NO 25219.
      (R,K) {20,20}
20: LSKEL KKKKKKKKKKKKKKKKKKKKKK KKKLG
      (K) {20}
21: SKELK KKKKKKKKKKKKKKKKKKKKKK KKLGD
      (K) {20}
22: KELKK KKKKKKKKKKKKKKKKKKKKKK KLGDE
      (K) {20}
23: ELKKK KKKKKKKKKKKKKKKKKKKKKK LGDEE
      (K) {20}

1  AA011342 ck: 7189 len: 106 ! Aa011342 Human polypeptide SEQ ID NO 25234.
      (R,K) {20,20}
61: FFKTK KKKKKKKKKKKKKKKKKKKKKK GGPLK
      (K) {20}

1  AA011346 ck: 7311 len: 85 ! Aa011346 Human polypeptide SEQ ID NO 25238.
      (R,K) {20,20}
62: ANHME KKKKKKKKKKKKKKKKKKKKKK SGGG
      (K) {20}

1  AA011352 ck: 1342 len: 51 ! Aa011352 Human polypeptide SEQ ID NO 25244.
      (R,K) {20,20}
25: SIFKK KKKKKKKKKKKKKKKKKKKKKK KNPPS
      (K) {20}
26: IFPKK KKKKKKKKKKKKKKKKKKKKKK NPPSL
      (K) {20}

1  AA011705 ck: 7660 len: 101 ! Aa011705 Human polypeptide SEQ ID NO 25597.
      (R,K) {20,20}
17: WAPLQ KKKKKKKKKKKKKKKKKKKKKK KKKGG
      (K) {20}
18: APLQK KKKKKKKKKKKKKKKKKKKKKK KKGGA
      (K) {20}
19: PLQKK KKKKKKKKKKKKKKKKKKKKKK KGGAP
      (K) {20}
20: LQKKK KKKKKKKKKKKKKKKKKKKKKK GGAPL
      (K) {20}

1  AA011799 ck: 8286 len: 39 ! Aa011799 Human polypeptide SEQ ID NO 25691.
      (R,K) {20,20}
7: NPVPS KKKKKKKKKKKKKKKKKKKKKK KKKKK
      (R,K) {20}
8: PPSVR KKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
9: PPSRK KKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}

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10: VSRKK KKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
11: SRKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
12: RKKKK KKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKGG
      (K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKGGG
      (K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKKKKK KGGGL
      (K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKKKKK GGGL

1  AA011820 ck: 8725 len: 42 ! Aa011820 Human polypeptide SEQ ID NO 25712.
      (R,K) {20,20}
17: NFKAL KKKKKKKKKKKKKKKKKKKKKK SSSLR
      (K) {20}

1  AA011828 ck: 4807 len: 41 ! Aa011828 Human polypeptide SEQ ID NO 25720.
      (R,K) {20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKKK KKKGG
      (K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKKK KRGGG
      (K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKKK RGGGF
      (K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKKK GGGRK
      (R,K) {20}

1  AA011831 ck: 1761 len: 29 ! Aa011831 Human polypeptide SEQ ID NO 25723.
      (R,K) {20,20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKA
      (K) {20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKKK KKKKG
      (K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKKK KKGAG
      (K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKKKKK KAGGG
      (K) {20}

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(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKK AGGA

AA011843 ck: 86 len: 36 1 Aa011843 Human polypeptide SEQ ID NO 25735.

1 (R, K) {20, 20}

(K) {20}
1: KKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
2: K KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
3: KK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
4: KKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
5: KKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
9: KKKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
10: KKKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
12: KKKKK KKKKKKKKKKKKKKKKK KKKSG

(K) {20}
13: KKKKK KKKKKKKKKKKKKKKKK KKS G

(K) {20}
14: KKKKK KKKKKKKKKKKKKKKKK KSG

(K) {20}
15: KKKKK KKKKKKKKKKKKKKKKK SG

AA011844 ck: 3574 len: 49 1 Aa011844 Human polypeptide SEQ ID NO 25736.

1 (R, K) {20, 20}

(K) {20}
18: ILYME KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
19: LYMEK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
20: YMEKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
21: MEKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}
22: EKKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}

23: KKKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}

24: KKKKK KKKKKKKKKKKKKKKKK KKKYS

(K) {20}

25: KKKKK KKKKKKKKKKKKKKKKK KKYSR

(K) {20}

26: KKKKK KKKKKKKKKKKKKKKKK KYSR

(K) {20}

27: KKKKK KKKKKKKKKKKKKKKKK YSR

AA011845 ck: 8375 len: 45 1 Aa011845 Human polypeptide SEQ ID NO 25737.

1 (R, K) {20, 20}

(K) {20}
15: SPSTL KKKKKKKKKKKKKKKKK KKKKK

(K) {20}

16: PSTLK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}

17: STLKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}

18: TLKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}

19: LKKKK KKKKKKKKKKKKKKKKK KKKKK

(K) {20}

20: KKKKK KKKKKKKKKKKKKKKKK KKKKT

(K) {20}

21: KKKKK KKKKKKKKKKKKKKKKK KKTG

(K) {20}

22: KKKKK KKKKKKKKKKKKKKKKK KKTG

(K) {20}

23: KKKKK KKKKKKKKKKKKKKKKK KTG

(K) {20}

24: KKKKK KKKKKKKKKKKKKKKKK TG

AA011849 ck: 284 len: 36 1 Aa011849 Human polypeptide SEQ ID NO 25741.

1 (R, K) {20, 20}

(K) {20}

13: KCLCE KKKKKKKKKKKKKKKKK KRKR

(K) {20}

14: CLCEK KKKKKKKKKKKKKKKKK RKR

(R, K) {20}

15: LCEKK KKKKKKKKKKKKKKKKK KR

(R, K) {20}

16: CEKKK KKKKKKKKKKKKKKKKK R

(R, K) {20}

17: EKKKK KKKKKKKKKKKKKKKKK KR

AA011886 ck: 5282 len: 97 1 Aa011886 Human polypeptide SEQ ID NO 25778.

1 (R, K) {20, 20}

(K) {20}

32: SNSSL KKKKKKKKKKKKKKKKK KKKKK

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33: NSSLK (K) [20]
34: SSLKK (K) [20]
35: SLKKK (K) [20]
36: LKKKK (K) [20]
37: KKKKK (K) [20]
38: KKKKK (K) [20]
39: KKKKK (K) [20]
40: KKKKK (K) [20]
41: KKKKK (K) [20]
42: KKKKK (K) [20]
43: KKKKK (K) [20]
44: KKKKK (K) [20]
45: KKKKK (K) [20]
46: KKKKK (K) [20]

AA011888 ck: 6488 len: 31 ! Aa011888 Human polypeptide SEQ ID NO 25780
(R,K) [20, 20]
2: D KKKKKKKKKKKKKKKKKKK KAGGG
(K) [20]
3: DK KKKKKKKKKKKKKKKKKKK KAGGGL

AA011903 ck: 8876 len: 48 ! Aa011903 Human polypeptide SEQ ID NO 25795
(R,K) [20, 20]
17: VIKYV KKKKKKKKKKKKKKKKKKK KKKKK
(K) [20]
18: IKYVK KKKKKKKKKKKKKKKKKKK KKKKK
(K) [20]
19: KYVKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) [20]
20: YVKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) [20]
21: VKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) [20]

```

1

22: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
23: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
24: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
25: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
26: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
27: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
28: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
29: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

AaO11999 ck: 3954 len: 47 i AaO11999 Human polypeptide SHQ ID NO 25891.

(R,K) {20,20}
(K) {20}
17: IDEGL KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
18: DEGLK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
19: EGLKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
20: GLKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
21: LKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
22: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
23: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
24: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20} GGAP

AaO12098 ck: 8434 len: 39 i AaO12098 Human polypeptide SHQ ID NO 25990.

(R,X) {20,20}
(K) {20}
13: TCKFG KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
14: CKFGK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
15: KCFGK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
16: FCGKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
17: GKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}
18: KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK
(K) {20}

19: KKKK (K) {20}
K
20: KKKK (K) {20}
K

AA012105 ck: 6255 len: 80 ! Aa012105 Human polypeptide SEQ ID NO 25997.
(R,K) {20,20}

53: EATSL KKKK (K) {20}
K
54: ATSLK KKKK (K) {20}
K
55: TSLK KKKK (K) {20}
K
56: SLKK KKKK (K) {20}
K
57: LKKK KKKK (K) {20}
K
58: KKKK KKKK (K) {20}
K

AA012179 ck: 1295 len: 69 ! Aa012179 Human polypeptide SEQ ID NO 26071.
(R,K) {20,20}

35: PRESV KKKK (K) {20}
K
36: KPSVK KKKK (K) {20}
K
37: FSVK KKKK (K) {20}
K
38: SVKK KKKK (K) {20}
K
39: VKKK KKKK (K) {20}
K
40: KKKK KKKK (K) {20}
K
41: KKKK KKKK (K) {20}
K
42: KKKK KKKK (K) {20}
K
43: KKKK KKKK (K) {20}
K
44: KKKK KKKK (K) {20}
K
45: KKKK KKKK (K) {20}
K
46: KKKK KKKK (K) {20}
K

AA012180 ck: 9840 len: 67 ! Aa012180 Human polypeptide SEQ ID NO 26072.
(R,K) {20,20}

38: CSYLP KKKK (K) {20}
K
39: SYLPK KKKK (K) {20}
K
40: YLPK KKKK (K) {20}
K
41: LPKK KKKK (K) {20}
K
42: PKKK KKKK (K) {20}
K
43: KKKK KKKK (K) {20}
K
44: KKKK KKKK (K) {20}
K
45: KKKK KKKK (K) {20}
K
46: KKKK KKKK (K) {20}
K
47: KKKK KKKK (K) {20}
K
48: KKKK KKKK (K) {20}
K

AA012187 ck: 4700 len: 60 ! Aa012187 Human polypeptide SEQ ID NO 26079.
(R,K) {20,20}

30: SCKL KKKK (K) {20}
K
31: CKLK KKKK (K) {20}
K
32: KCLK KKKK (K) {20}
K
33: KKKK KKKK (K) {20}
K
34: LKKK KKKK (K) {20}
K
35: KKKK KKKK (K) {20}
K
36: KKKK KKKK (K) {20}
K
37: KKKK KKKK (K) {20}
K
38: KKKK KKKK (K) {20}
K
39: KKKK KKKK (K) {20}
K
40: KKKK KKKK (K) {20}
K
41: KKKK KKKK (K) {20}
K

1 AAO12203 ck: 4083 len: 41 ! Aao12203 Human polypeptide SEQ ID NO 26095.
(R, K) {20, 20}
(K) {20}

6: SWCCL KKKKKKKKKKKKKKKKKKKKK KKPFG

7: WCCLK KKKKKKKKKKKKKKKKKKKKK KKPFG
(K) {20}

8: CCLKK KKKKKKKKKKKKKKKKKKKKK KPPGG
(K) {20}

9: CLKKK KKKKKKKKKKKKKKKKKKKKK PGGGA
(K) {20}

AAO12215 ck: 6903 len: 73 ! Aao12215 Human polypeptide SEQ ID NO 26107.

(R, K) {20, 20}

(K) {20}

50: PPFLP KKKKKKKKKKKKKKKKKKKKK KTGCG

51: PFLPK KKKKKKKKKKKKKKKKKKKKK TGG
(K) {20}

AAO12243 ck: 8474 len: 39 ! Aao12243 Human polypeptide SEQ ID NO 26135.

(R, K) {20, 20}

(K) {20}

11: MISPI KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

12: ISPIK KKKKKKKKKKKKKKKKKKKKK KKKKG

(K) {20}

13: SPIKK KKKKKKKKKKKKKKKKKKKKK KKKGG

(K) {20}

14: FIYKK KKKKKKKKKKKKKKKKKKKKK KKGGP

(K) {20}

15: IKKKK KKKKKKKKKKKKKKKKKKKKK KGGPL

(K) {20}

16: KKKKK KKKKKKKKKKKKKKKKKKKKK GGPL

AAO12250 ck: 1538 len: 69 ! Aao12250 Human polypeptide SEQ ID NO 26142.

(R, K) {20, 20}

(K) {20}

31: SPSNL KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

32: PSNLK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

33: SNLKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

34: NLKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

35: LKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

36: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

37: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

1 38: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

39: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

40: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

41: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKRG
(K) {20}

42: KKKKK KKKKKKKKKKKKKKKKKKKKK KRRGG
(K) {20}

43: KKKKK KKKKKKKKKKKKKKKKKKKKK KRGCG
(K) {20}

44: KKKKK KKKKKKKKKKKKKKKKKKKKK RGGGA
(K) {20}

45: KKKKK KKKKKKKKKKKKKKKKKKKKK GGGAF
(R, K) {20}

AAO12274 ck: 2739 len: 37 ! Aao12274 Human polypeptide SEQ ID NO 26166.

(R, K) {20, 20}

(K) {20}

14: KGLIN KKKKKKKKKKKKKKKKKKKKK KKKGG
(K) {20}

15: GLINK KKKKKKKKKKKKKKKKKKKKK KKG
(K) {20}

16: LINKK KKKKKKKKKKKKKKKKKKKKK KG
(K) {20}

17: INKKK KKKKKKKKKKKKKKKKKKKKK G

1 AAO12280 ck: 7481 len: 66 ! Aao12280 Human polypeptide SEQ ID NO 26172.

(R, K) {20, 20}

(K) {20}

29: HTPPL KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

30: IPPLK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

31: PPLKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

32: PLKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

33: LKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

34: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

35: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

36: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

37: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}

38: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKS

1
39: KKKKK (K) {20}
40: KKKKK (K) {20}
41: KKKKK (K) {20}
42: KKKKK (K) {20}
AA012447 ck: 5017 len: 47 | Aa012447 Human polypeptide SEQ ID NO 26339.
(R, K) {20, 20}
(K) {20}
14: HLVCE KKKKKKKKKKKKKKKKKKKR KRGA
(K) {20}
15: LVCEK KKKKKKKKKKKKKKKKKKKR RGGAL
(R, K) {20}
16: VCEKK KKKKKKKKKKKKKKKKKKKR GGALK
AA012476 ck: 837 len: 104 | Aa012476 Human polypeptide SEQ ID NO 26368.
(R, K) {20, 20}
(K) {20}
65: RFCHQ KKKKKKKKKKKKKKKKKKKR KGFFP
(K) {20}
66: FCHQK KKKKKKKKKKKKKKKKKKKR GPPFW
AA012548 ck: 6973 len: 74 | Aa012548 Human polypeptide SEQ ID NO 26440.
(R, K) {20, 20}
(K) {20}
38: AVLPL KKKKKKKKKKKKKKKKKKKR KKGCG
(K) {20}
39: VLPLK KKKKKKKKKKKKKKKKKKKR KGGGV
(K) {20}
40: LPLKK KKKKKKKKKKKKKKKKKKKR GGGVF
AA012553 ck: 1903 len: 33 | Aa012553 Human polypeptide SEQ ID NO 26445.
(R, K) {20, 20}
(K) {20}
9: SCCFI KKKKKKKKKKKKKKKKKKKR KRGAP
(K) {20}
10: CCFIK KKKKKKKKKKKKKKKKKKKR RGAP
(R, K) {20}
11: CFIKK KKKKKKKKKKKKKKKKKKKR GAP
AA013164 ck: 2798 len: 71 | Aa013164 Human polypeptide SEQ ID NO 27056.
(R, K) {20, 20}
(R, K) {20}
35: RPPLX RKKKKKKKKKKKKKKKKKKR KEMFK
(K) {20}
36: PPLXR KKKKKKKKKKKKKKKKKKKR EMFKR
AA013576 ck: 4846 len: 99 | Aa013576 Human polypeptide SEQ ID NO 27468.

1
53: TNNLI (R, K) {20, 20}
(K) {20}
54: NNLIK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
55: NLIK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
56: LIKK KKKKKKKKKKKKKKKKKKKR KKKKG
(K) {20}
57: IKKK KKKKKKKKKKKKKKKKKKKR KKKGG
(K) {20}
58: KKKKK KKKKKKKKKKKKKKKKKKKR KKGCG
(K) {20}
59: KKKKK KKKKKKKKKKKKKKKKKKKR KGGGP
(K) {20}
60: KKKKK KKKKKKKKKKKKKKKKKKKR GGPP
AA013785 ck: 6241 len: 100 | Aa013785 Human polypeptide SEQ ID NO 27677.
(R, K) {20, 20}
(K) {20}
41: LYAPP KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
42: YAPPK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
43: APPKK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
44: PPKKK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
45: PKKKK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
46: KKKKK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
47: KKKKK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
48: KKKKK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
49: KKKKK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
50: KKKKK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
51: KKKKK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
52: KKKKK KKKKKKKKKKKKKKKKKKKR KKKKK
(K) {20}
53: KKKKK KKKKKKKKKKKKKKKKKKKR KKKKA
(K) {20}
54: KKKKK KKKKKKKKKKKKKKKKKKKR KKKAG
(K) {20}

55: KKKKK KKKKKKKKKKKKKKKKKKKKK KTAGG
(K) {20}
56: KKKKK KKKKKKKKKKKKKKKKKKKKK KAGGG
(K) {20}
57: KKKKK KKKKKKKKKKKKKKKKKKKKK AGGGG
AAU17983 ck: 7016 len: 315 ! Aau17983 Human immunoglobulin polypeptide S
(R, K) {20, 20}
(R, K) {20}
273: QVFAP RKKKKKKKKKKKKKKKKKKKK KGGRS
(K) {20}
274: VFAPR KKKKKKKKKKKKKKKKKKKKK GGRSR
AAU18049 ck: 7611 len: 315 ! Aau18049 Human immunoglobulin polypeptide S
(R, K) {20, 20}
(R, K) {20}
273: QVFAP RKKKKKKKKKKKKKKKKKKKK KGGRS
(K) {20}
274: VFAPR KKKKKKKKKKKKKKKKKKKKK GGRSR
AAM14119 ck: 5383 len: 86 ! Aam14119 Peptide #553 encoded by probe for
(R, K) {20, 20}
(R, K) {20}
15: RRRRG RRRRRKKKKKKKKKKKKKK KKRK
(R, K) {20}
16: RRRGR RRRKKKKKKKKKKKKKKKK KRRR
(R, K) {20}
17: RRGRR RRRKKKKKKKKKKKKKKKK KRRR
(R, K) {20}
18: RGRRR RKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
19: GRRRR RKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
20: RRRRR KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
21: RRRRK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
22: RRRKK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
23: RRRKK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
24: RKKKK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
25: KKKKK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
26: KKKKK RKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
27: KKKKK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
28: KKKKK KKKKKKKKKKKKKKKKKKKR RRRR

29: KKRKK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
30: KKKKK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
31: KKKKK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
32: KKKKK RKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
33: KKKKK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
34: KKKKK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
35: KKKKK KKKKKKKKKKKKKKKKKKKR RRRR
(R, K) {20}
36: KKKKK KRRRRKKKKKKKKKKKKKKR RRRR
(R) {20}
37: RKKKK RRRRRKKKKKKKKKKKKKKR RRRR
(R) {20}
38: KKKKK RRRRRKKKKKKKKKKKKKKR RRRR
(R) {20}
39: KKKKK RRRRRKKKKKKKKKKKKKKR RRRR
(R) {20}
40: KKKRR RRRRRKKKKKKKKKKKKKKR RRRR
(R) {20}
41: KRRRR RRRRRKKKKKKKKKKKKKKR RRRR
(R) {20}
42: RRRRR RRRRRKKKKKKKKKKKKKKR RRRR
(R) {20}
43: RRRRR RRRRRKKKKKKKKKKKKKKR RRRR
(R) {20}
44: RRRRR RRRRRKKKKKKKKKKKKKKR RRRR
(R) {20}
45: RRRRR RRRRRKKKKKKKKKKKKKKR RRRR
(R) {20}
AAM14961 ck: 1334 len: 86 ! Aam14961 Peptide #1395 encoded by probe for
(R, K) {20, 20}
(R, K) {20}
57: EEEEG RRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}
58: EEEGR RRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}
59: EEEGR RRRKKKKKKKKKKKKKKKK KKKK
(R, K) {20}
60: EGGRR KKKKKKKKKKKKKKKKKKKK KKKK
(R, K) {20}
61: GRRRK KKKKKKKKKKKKKKKKKKKK KKKK
(R, K) {20}
62: RRRKK KKKKKKKKKKKKKKKKKKKK KKKK

[illegible]

AAM15038 ck: 9082 len: 167 ! Aam15038 Peptide #1472 encoded by probe for

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(R,K) {20,20}
(R,K) {20}
33: EGGG RRRRRRRRRRRRRRRRR RGGG

```

34: EGRGR RRRRRKKRRKKRRKKRR (R, K) { 20 } RGGGR

35: $\text{GGR} \text{RRRRKKRRKKRRKKRR} \text{GGGR}$
 $(R, K) \{20\}$

AAM15527 ck: 2276 len: 89 ! Aam15527 Peptide #1961 encoded by probe for

$$\begin{array}{l} (R, K) \{20, 20\} \\ (R, K) \{20\} \end{array}$$
$$(K) \{20\}$$
$$(K) \{20\}$$
$$(R, K) \{20\}$$

(R, K) {20}

AAM15826 ck: 1939 len: 130 ! Aam15826 Peptide #2260 encoded by probe for

1

$$\begin{aligned} & (\mathbb{R}, \kappa) \{20, 20\} \\ & (\mathbb{R}, \kappa) \{20\} \end{aligned}$$

43: GRKER RRRRRRRRRRKRK RR

44: RKER RRRRRRRRRRRR RRRR

45: KERR RRRRRRRRRRRRRRRRR RRRK

46: E R R R R R R R R R R R R R K K K K K

47: RRRR RRRRRRRRRRRR RRRR

48: RRRR RRRRRRRRRKKRRRRR KKKKK

49: RRRRR (R,K){20} RRRRRRRRRKKRRRRRRK KKKKK

```
50: RRRR RRRRRRRKKKKRRRRKK KKKK
      (R,K){20}
```

```
51: RRRR RRRRRRKKRRRRKKK KKKK
      (R,K){20}
```

```

(R,X){20}
52: RRRR RRRRRKKRRRRKKKK KEEE

```

53: RRRR RRRRKKRRRRKKKK KEEE (R,K){20}

```

54: RRRRR RRRRRKKRRRRRRKKKKK EEEEE
      (R,K) { 20 }

```

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AAm16123 ck: 3607 len: 88 ! Aam16123 Peptide #2557 encoded by probe for
```

39. ERKRE KKKKKKKKKKKKKKKKKKKKKKKKKKK
 (R,K) { 20, 20 }
 (K) { 20 }

40: RPREX KKKKKKKKKKKKKKKKKKKKKKKKKKKKK

```

41: KREKK XXXXXXXXXXXXXXXXXXXX XXXXX
      (K) { 20 }

```

(K) { 20 }.

REKK KKKKKKKKKKKKKKKKK

```

      (X) { 20 }
43: EKKK KKKKKKKKKKKKKKKKKKK KKKK

```

44: KKKK KKKKKKKKKKKKKKKKK KKKK
 (K) { 20 }

45: KKKKK (K) { 20 } KKKKK

46: KKKK (K) { 20 } KKKK

47: KKKK (k) {20} KKKK

48: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

49: KKKK KKKKKKKKKKKKKKKKK KKKK
 (K) { 20 }

(K) { 20 }

50 : KKKKK KKKKKKKKKKKKKKKKKKKKKKKKKKK

(K) { 20 }

51: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) { 20 }

52 : KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) { 20 }

53 : KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) { 20 }

54 : KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

(K) { 20 }

55 : KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

56: KKKK (K) {20}
57: KKKK (K) {20}
58: KKKK (K) {20}
59: KKKK (K) {20}
60: KKKK (K) {20}
61: KKKK (K) {20}
62: KKKK (K) {20}
63: KKKK (K) {20}
64: KKKK (K) {20}
65: KKKK (K) {20}
66: KKKK (K) {20}
67: KKKK (K) {20}
68: KKKK (K) {20}
69: KKKK (K) {20}

AAm1691 ck: 3937 len: 85 ! Aam1691 Peptide #3125 encoded by probe for

1
(R, K) {20, 20}
1: KKKK (K) {20}
2: K KKKK (K) {20}
3: KK KKKK (K) {20}
4: KK KKKK (K) {20}
5: KKKK (K) {20}
6: KKKK (K) {20}
7: KKKK (K) {20}
8: KKKK (K) {20}
9: KKKK (K) {20}

10: KKKK (K) {20}
11: KKKK (K) {20}

AAm1738 ck: 2686 len: 71 ! Aam1738 Peptide #3822 encoded by probe for

1
(R, K) {20, 20}
20: KKKK (K) {20}
21: KKKK (K) {20}
22: KKKK (K) {20}
23: KKKK (K) {20}
24: KKKK (K) {20}
25: KKKK (R, K) {20}
26: KKKK (R, K) {20}
27: KKKK (R, K) {20}
28: KKKK (R, K) {20}
29: KKKK (R, K) {20}
30: KKKK (R, K) {20}
31: KKKK (R, K) {20}
32: KKKK (R, K) {20}
33: KKKK (R, K) {20}
34: KKKK (R, K) {20}
35: KKKK (R, K) {20}
36: KKKK (R, K) {20}
37: KKKK (R, K) {20}
38: KKKK (R, K) {20}
39: KKKK (R, K) {20}
40: KKKK (R, K) {20}

1

```
43: GRKER RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
44: RKERR RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
45: KEERRR RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
46: ERRRRR RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
47: RRRRRR RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
48: RRRRRR RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
49: RRRRRR RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
50: RRRRRR RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
51: RRRRRR RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
52: RRRRRR RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
53: RRRRRR RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
54: RRRRRR RRRRRRRRRRRRRRRRRRRR RRRRR (R,K){20}
AAM28616 ck: 3607 len: 88 ! Aam28616 Peptide #2653 encoded by probe for
(R,K){20,20}
(R,K){20}
39: ERKRE KKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20,20}
40: RKREK KKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20}
41: KREKK KKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20}
42: REKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20}
43: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20}
44: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20}
45: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20}
46: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20}
47: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20}
48: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20}
49: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20}
```

1

```
50: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
51: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
52: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
53: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
54: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
55: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
56: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
57: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
58: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
59: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
60: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
61: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
62: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
63: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
64: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
65: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
66: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
67: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
68: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
69: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
AAM29178 ck: 3937 len: 85 ! Aam29178 Peptide #3215 encoded by probe for
(R,K){20,20}
(R,K){20}
1: KKKKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK (R,K){20,20}
2: K KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
3: KK KKKKKKKKKKKKKKKKKKKKKKKKKKK KKKKK (K){20}
```


3: RR RRRRRRRRRRRRRRRR RT
(R,K){20}
4: RRR RRRRRRRRRRRRRRRR T
(R,K){20}

AAM37794 ck: 3301 len: 52 i Aam37794 Peptide #11831 encoded by probe fd

12: KKKKN KKKKKKKKKKKKKKKK KKKK
(R,K){20,20}
13: KKKKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
14: KKNKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
15: KNKKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
16: NKKKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
17: KKKKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
18: KKKKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
19: KKKKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
20: KKKKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
21: KKKKK RKKKKKKKKKKKKKKK KKKK
(R,K){20}
22: KKKKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
23: KKKKK RKKKKKKKKKKKKKKK KKKK
(R,K){20}
24: KKKKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
25: KKKKK RKKKKKKKKKKKKKKK KKKK
(R,K){20}
26: RKKKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
27: KKKKK RKKKKKKKKKKKKKKK KKKK
(R,K){20}
28: RKKKK KKKKKKKKKKKKKKKK KKKK
(R,K){20}
29: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
30: RKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
31: KKKKK KKKKKKKKKKKKKKKK AF
(K){20}

AAM38273 ck: 4895 len: 51 i Aam38273 Peptide #12310 encoded by probe fd

18: LFKPM RKKRRKKRRKKKKKKRR KKLTT
(R,K){20,20}
19: FKPMR KRRKKKKRRKKKKRRK KLTTT
(R,K){20}
20: KPMRK RRRKKRRKKKKKKRRK LTTT
(R,K){20}

AAU04283 ck: 8137 len: 45 i Aau04283 Trimeric fusogenic peptide #2 used

4: YKA KKKKKKKKKKKKKKKKK KKKK
(R,K){20,20}
5: YKAK KKKKKKKKKKKKKKKK KKKK
(K){20}
6: YKAKK KKKKKKKKKKKKKKKK KKKK
(K){20}
7: KAKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
8: AKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
9: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
10: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
11: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
12: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
13: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
14: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
15: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
16: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
17: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
18: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
19: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
20: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
21: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}
22: KKKKK KKKKKKKKKKKKKKKK KKKK
(K){20}

23: KKKKK KKKKKKKKKKKKKKKKK KWK
24: KKKKK KKKKKKKKKKKKKKKKK KWK
AaU04285 ck: 4361 len: 59 ! AaU04285 Nuclear ligand #2 used in nucleic
(R, K) {20, 20}
(K) {20}

18: APyKA KKKKKKKKKKKKKKKKK KKKKK

19: PYKAK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

20: YKAKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

21: KAKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

22: AKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

23: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

24: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

25: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

26: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

27: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

28: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

29: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

30: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

31: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

32: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

33: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

34: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

35: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

36: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

37: KKKKK KKKKKKKKKKKKKKKKK KWK
(K) {20}

38: KKKKK KKKKKKKKKKKKKKKKK KWK
(K) {20}

1

AaU04287 ck: 4925 len: 100 ! AaU04287 Poly-L-Lysine used in nucleic acid
(R, K) {20, 20}
(K) {20}

1: KKKKKKKKKKKKKKKKKKK KKKKK

2: K KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

3: KK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

4: KKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

5: KKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

6: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

7: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

8: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

9: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

10: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

11: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

12: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

13: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

14: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

15: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

16: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

17: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

18: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

19: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

20: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

21: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

22: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

23: KKKKK KKKKKKKKKKKKKKKKK KKKKK
(K) {20}

73: KKKKK (K) {20} KKKKK
 74: KKKKK (K) {20} KKKKK
 75: KKKKK (K) {20} KKKKK
 76: KKKKK (K) {20} KKKKK
 77: KKKKK (K) {20} KKKKK
 78: KKKKK (K) {20} KKKKK
 79: KKKKK (K) {20} KKKKK
 80: KKKKK (K) {20} KKKKK
 81: KKKKK (K) {20} KKKKK

AAW01857 ck: 5383 len: 86 i Aaw01857 Peptide #539 encoded by probe for

1

15: RRRRG (R,K) {20,20} KKKKK
 16: RRRGR (R,K) {20} KRRRR
 17: RRGRR (R,K) {20} RRRRR
 18: RGRRR (R,K) {20} RRRRR
 19: GRRRR (R,K) {20} RRRRR
 20: RRRRR (R,K) {20} RRRRR
 21: RRRRK (R,K) {20} RRRRR
 22: RRRKK (R,K) {20} RRRRR
 23: RRRKK (R,K) {20} RRRRR
 24: RKKKK (R,K) {20} RRRRR
 25: KKKKK (R,K) {20} RRRRR
 26: KKKKK (R,K) {20} RRRRR
 27: KKKKK (R,K) {20} RRRRR
 28: KKKKK (R,K) {20} RRRRR

1

29: KKKKK (R,K) {20} RRRRR
 30: KKKKK (R,K) {20} RRRRR
 31: KKKKK (R,K) {20} RRRRR
 32: KKKKK (R,K) {20} RRRRR
 33: KKKKK (R,K) {20} RRRRR
 34: KKKKK (R,K) {20} RRRRR
 35: KKKKK (R,K) {20} RRRRR
 36: KKKKK (R,K) {20} RRRRR
 37: KKKKK (R) {20} RRRRR
 38: KKKKK (R) {20} RRRRR
 39: KKKKK (R) {20} RRRRR
 40: KKKKK (R) {20} RRRRR
 41: KKKKK (R) {20} RRRRR
 42: KKKKK (R) {20} RRRRR
 43: KKKKK (R) {20} RRRRR
 44: KKKKK (R) {20} RRRRR
 45: KKKKK (R) {20} RRRRR
 46: KKKKK (R) {20} RRRRR
 47: KKKKK (R) {20} RRRRR
 48: KKKKK (R) {20} RRRRR
 49: KKKKK (R) {20} RRRRR
 50: KKKKK (R) {20} RRRRR
 51: KKKKK (R) {20} RRRRR
 52: KKKKK (R) {20} RRRRR
 53: KKKKK (R) {20} RRRRR
 54: KKKKK (R) {20} RRRRR
 55: KKKKK (R) {20} RRRRR
 56: KKKKK (R) {20} RRRRR
 57: KKKKK (R) {20} RRRRR
 58: KKKKK (R) {20} RRRRR
 59: KKKKK (R) {20} RRRRR
 60: KKKKK (R) {20} RRRRR
 61: KKKKK (R) {20} RRRRR
 62: KKKKK (R) {20} RRRRR

AAW02687 ck: 1334 len: 86 i Aaw02687 Peptide #1369 encoded by probe for

(R,K){20}
63: RRRKK KRRKKKKKKKKKKKKKKKKKK
(R,K){20}
64: RKKKK RRRKKKKKKKKKKKKKKKKKK
(R,K){20}
65: KKKKK KRRKKKKKKKKKKKKKKKKKK
(R,K){20}
66: KKKKK RKKKKKKKKKKKKKKKKKK K
(K){20}
67: KKKKK KKKKKKKKKKKKKKKKKKK

1
AAM02768 ck: 9082 len: 167 1 Aam02768 Peptide #1450 encoded by probe for
(R,K){20,20}
33: EEEGG RRRRRRRRRRRRRRRRRRRR RRGCG
(R,K){20}
34: EGGGR RRRRRRRRRRRRRRRRRRRR RGGGR
(R,K){20}
35: GGGRR RRRRRRRRRRRRRRRRRRRR GGGRR

1
AAM03278 ck: 2276 len: 89 1 Aam03278 Peptide #1960 encoded by probe for
(R,K){20,20}
23: EEEEE KKKKKRRRRRRRRRRRRRRR EEEKK
(R,K){20}
46: KKEEE KKKKKKKKKKKKKKKKKKK KKEEE
(K){20}
47: KEEKK KKKKKKKKKKKKKKKKKKK RKEEE
(R,K){20}
48: EEEKK KKKKKKKKKKKKKKKKKKK KEEEE
(R,K){20}
49: EEEKK KKKKKKKKKKKKKKKKKKK EEEEE

1
AAM03564 ck: 1939 len: 130 1 Aam03564 Peptide #2246 encoded by probe for
(R,K){20,20}
42: EGGKE RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}
43: GKKER RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}
44: RKERR RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}
45: KERRR RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}
46: ERRRR RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}
47: RRRRR RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}
48: RRRRR RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}

49: RRRRR RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}
50: RRRRR RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}
51: RRRRR RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}
52: RRRRR RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}
53: RRRRR RRRRRRRRRRRRRRRRRRRR RRRRR
(R,K){20}
54: RRRRR RRRRRRRRRRRRRRRRRRRR RRRRR

1
AAM03848 ck: 3607 len: 88 1 Aam03848 Peptide #2530 encoded by probe for
(R,K){20,20}
39: EKKKE KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
40: RKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
41: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
42: RKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
43: EKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
44: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
45: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
46: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
47: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
48: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
49: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
50: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
51: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
52: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
53: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
54: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}
55: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K){20}

56: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
57: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
58: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
59: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
60: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
61: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
62: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
63: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
64: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
65: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
66: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
67: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
68: KKKKK KKKKKKKKKKKKKKKKKKKKK K
(K) {20}
69: KKKKK KKKKKKKKKKKKKKKKKKKKK

1
AAM04408 ck: 3937 len: 85 ! Aam04408 Peptide #3090 encoded by probe for

(R, K) {20, 20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
2: K KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
3: KK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
4: KKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
5: KKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
6: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
7: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
8: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

10: KKKKK KKKKKKKKKKKKKKKKKKKKK KEEEE
(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKKKK BEEEE

1
AAM06100 ck: 1560 len: 88 ! Aam06100 Peptide #4782 encoded by probe for

(R, K) {20, 20}
43: RRRRG RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
44: RRRGR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
45: RRGRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
46: RGRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
47: GRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
48: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
49: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
50: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
51: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
52: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
53: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
54: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
55: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
56: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
57: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
58: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
59: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
60: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
61: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}
62: RRRRR RRRRRRRRRRRRRRRRRRRRR RRRRR
(R) {20}

1
AAG73687 ck: 3063 len: 29 ! Aag73687 Human colon cancer antigen protein
(R, K) {20, 20}

(K) {20}
8: MMTTF KKKKKKKKKKKKKKKKKKKKK KX
(K) {20}
9: MMTFK KKKKKKKKKKKKKKKKKKKKK X

1 AAG73729 ck: 783 len: 83 ! Aag73729 Human colon cancer antigen protein

(R,K) {20,20}
(K) {20}

49: LGPCE KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
50: GPCEK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
51: PCEKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
52: CEKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
53: EKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
54: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
55: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

56: KKKKK KKKKKKKKKKKKKKKKKKKKK GGRKK
AAG73810 ck: 3374 len: 88 ! Aag73810 Human colon cancer antigen protein

(R,K) {20,20}
(K) {20}

44: FGQTX KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
45: GQTXK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
46: QTXKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
47: TXKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
48: XKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
49: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
50: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
51: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
52: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
53: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
54: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
55: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

(K) {20}
56: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
57: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

58: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

59: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

60: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

61: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

62: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

63: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

64: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}

65: KKKKK KKKKKKKKKKKKKKKKKKKKK GGPX
(K) {20}

1 AAG73895 ck: 1887 len: 43 ! Aag73895 Human colon cancer antigen protein

(R,K) {20,20}
(R,K) {20}

18: VRPRV RKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
19: RPRVR KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
20: PRVRK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
21: RVRKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
22: VRKKK KKKKKKKKKKKKKKKKKKKKK GG

AAG74218 ck: 8659 len: 104 ! Aag74218 Human colon cancer antigen protein

(R,K) {20,20}
(K) {20}

75: PLGGQ KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
76: LGGQK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
77: GGQKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
78: GQKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
79: QKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
80: KKKKK KKKKKKKKKKKKKKKKKKKKK KKKKK

1

AAG74527 ck: 2664 len: 40 i Aag74527 Human colon cancer antigen protein
(R,K){20,20}
(K){20}
9: CULLY KKKKKKKKKKKKKKKKKKK KKKK
10: LLLYK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
11: LLYK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
12: LYKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
13: YKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
14: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
15: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
16: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
17: KKKK KKKKKKKKKKKKKKKKKKK KKKK

1

AAG74650 ck: 1596 len: 69 i Aag74650 Human colon cancer antigen protein
(R,K){20,20}
(K){20}
36: LQCRQ KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
37: QCRQK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
38: CROK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
39: RQKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
40: QKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
41: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
42: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
43: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
44: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
45: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
46: KKKK KKKKKKKKKKKKKKKKKKK KKKK

1

AAG74793 ck: 8497 len: 152 i Aag74793 Human colon cancer antigen protein
(R,K){20,20}
(K){20}
122: SHTQ KKKKKKKKKKKKKKKKKKK KKKK

1

123: SHTQ KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
AAG74907 ck: 1215 len: 98 i Aag74907 Human colon cancer antigen protein
(R,K){20,20}
(K){20}
57: NLRKE KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
58: LRKEK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
59: RKEKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
60: KEKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
61: EKKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
62: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
63: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
64: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
65: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
66: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
67: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
68: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
69: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
70: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
71: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
72: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
73: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
74: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
75: KKKK KKKKKKKKKKKKKKKKKKK KKKK
(K){20}
76: KKKK KKKKKKKKKKKKKKKKKKK KKKK

1

AAG75215 ck: 3913 len: 155 i Aag75215 Human colon cancer antigen protein
(R,K){20,20}
(K){20}

135: RSSAP KKKKKKKKKKKKKKKKKKK K
(K) {20}
136: SSAPK KKKKKKKKKKKKKKKKKKK

1
AAG75886 ck: 4235 len: 71 1 Aag75886 Human colon cancer antigen protein
(R, K) {20, 20}
(K) {20}
47: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
48: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

1
AAE01796 ck: 4416 len: 72 1 Aae01796 Human gene 27 encoded secreted pro
(R, K) {20, 20}
(K) {20}
47: LPTFL KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
48: PTFLK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
49: TPLK KKKKKKKKKKKKKKKKKKK KKKK
(K) {20}
50: FLKK KKKKKKKKKKKKKKKKKKK KKK
(K) {20}
51: LKKKK KKKKKKKKKKKKKKKKKKK KK
(K) {20}
52: KKKKK KKKKKKKKKKKKKKKKKKK K
(K) {20}
53: KKKKK KKKKKKKKKKKKKKKKKKK

1
AAE01848 ck: 5584 len: 73 1 Aae01848 Human gene 27 encoded secreted pro
(R, K) {20, 20}
(K) {20}
47: LPTFL KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
48: PTFLK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
49: TPLK KKKKKKKKKKKKKKKKKKK KKKKI
(K) {20}
50: FLKK KKKKKKKKKKKKKKKKKKK KKKI
(K) {20}
51: LKKKK KKKKKKKKKKKKKKKKKKK KKI
(K) {20}
52: KKKKK KKKKKKKKKKKKKKKKKKK KI
(K) {20}
53: KKKKK KKKKKKKKKKKKKKKKKKK I

1
AAB90574 ck: 1431 len: 530 1 Aab90574 Human secreted protein, SEQ ID NO:
(R, K) {20, 20}
(K) {20}
511: LHAP KKKKKKKKKKKKKKKKKKK

1
AAB45846 ck: 8137 len: 45 1 Aab45846 Nucleic acid transporter system pei
(R, K) {20, 20}
(K) {20}

4: YKA KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
5: YKAK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
6: YKAK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
7: KAKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
8: AKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
9: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
10: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
11: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
12: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
13: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
14: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
15: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
16: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
17: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
18: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
19: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
20: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
21: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
22: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
23: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
(K) {20}
24: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK

1
AAB45848 ck: 4361 len: 59 1 Aab45848 Nucleic acid transporter system pei
(R, K) {20, 20}
(K) {20}
18: APYKA KKKKKKKKKKKKKKKKKKK KKKKK


```

19: PYAAK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
20: YKAAK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
21: KAAKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
22: AKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
23: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
24: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
25: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
26: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
27: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
28: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
29: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
30: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
31: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
32: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
33: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
34: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
35: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
36: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
37: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
38: KKKKK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}

AAB45850 ck: 4925 len: 100 ! Aab45850 Nucleic acid transporter system pe
      (R,K) {20,20}
1: KKKKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
2: K KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}
3: KK KKKKKKKKKKKKKKKKKKK KKKKK
      (K) {20}

```

[illegible]

77: KKKK (K) {20} KKKK
78: KKKK (K) {20} KKKK
79: KKKK (K) {20} KKKK
80: KKKK (K) {20} K
81: KKKK (K) {20} KKKK

AAB50247 ck: 8085 len: 154 ! Aab50247 Human breast cancer associated B72

(R, K) {20, 20}

(K) {20}

114: TOLRQ KKKK KKKK KKKK KKKK KKKK

(K) {20}

115: QLRQK KKKK KKKK KKKK KKKK KKKK

(K) {20}

116: LRQK KKKK KKKK KKKK KKKK KKKK

(K) {20}

117: RQK KKKK KKKK KKKK KKKK KKKK

(K) {20}

118: QKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

119: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

120: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

121: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

122: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

123: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

124: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

125: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

126: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

127: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

128: KKKK KKKK KKKK KKKK KKKK KKKK

ABJ37738 ck: 8085 len: 154 ! Abj37738 Human tumour-related protein - SEQ

(R, K) {20, 20}

(K) {20}

114: TOLRQ KKKK KKKK KKKK KKKK KKKK

(K) {20}

115: QLRQK KKKK KKKK KKKK KKKK KKKK

1

116: LRQK KKKK KKKK KKKK KKKK KKKK
117: RQK KKKK KKKK KKKK KKKK KKKK
118: QKKK KKKK KKKK KKKK KKKK KKKK
119: KKKK (K) {20} KKKK
120: KKKK (K) {20} KKKK
121: KKKK (K) {20} KKKK
122: KKKK (K) {20} KKKK
123: KKKK (K) {20} KKKK
124: KKKK (K) {20} KKKK
125: KKKK (K) {20} KKKK
126: KKKK (K) {20} KKKK
127: KKKK (K) {20} KKKK
128: KKKK (K) {20} KKKK

ABR00951 ck: 6732 len: 139 ! ABR00951 Human gene 5-encoded secreted prot

(R, K) {20, 20}

(K) {20}

91: HFRPG KKKK KKKK KKKK KKKK KKKK

(K) {20}

92: FRPG KKKK KKKK KKKK KKKK KKKK

(K) {20}

93: RPGK KKKK KKKK KKKK KKKK KKKK

(K) {20}

94: PGKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

95: GKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

96: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

97: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

98: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

99: KKKK KKKK KKKK KKKK KKKK KKKK

(K) {20}

100: KKKK KKKK KKKK KKKK KKKK KKKK

101: KKKKK (K) {20} KKKKK
 102: KKKKK (K) {20} KKKKK
 103: KKKKK (K) {20} KKKKK
 104: KKKKK (K) {20} KKKKK
 105: KKKKK (K) {20} KKKKK
 106: KKKKK (K) {20} KKKKK
 107: KKKKK (K) {20} KKKKK
 108: KKKKK (K) {20} KKKKK
 109: KKKKK (K) {20} KKKKK
 110: KKKKK (K) {20} KKKKK
 111: KKKKK (K) {20} KKKKK
 112: KKKKK (K) {20} KKKKK
 113: KKKKK (K) {20} KKKKK
 114: KKKKK (K) {20} KKKKK
 115: KKKKK (K) {20} KKKKK
 116: KKKKK (K) {20} KKKKK
 117: KKKKK (K) {20} KKKKK
 118: KKKKK (K) {20} KKKKK
 119: KKKKK (K) {20} K
 120: KKKKK (K) {20} KKKKK

ABR01117 ck: 9531 len: 66 ! ABR01117 Human gene 171-encoded secreted pr

41: MMTVX (R,K) {20,20} KKKKK
 42: WTVXK (K) {20} KKKKK
 43: TVXKK (K) {20} KKKKK

44: VXXXX (K) {20} KKKKK
 45: XXXXX (K) {20} KKKKK
 46: KKKKK (K) {20} KKKKK
 47: KKKKK (K) {20} KKKKK

AAE33919 ck: 185 len: 22 ! Aae33919 Secretion domain peptide #17. 5/20

1: (R,K) {20,20} KKKKK
 1: RRRRRRRRRRRRRRRRRR GC

ABG73150 ck: 5536 len: 84 ! Abg73150 Single-chain antigen-binding (eFv)

1: (R,K) {20,20} KKKKK
 1: KKKKKKKKKKKKKKKKKKK KKKKK
 2: K KKKKKKKKKKKKKKKKK KKKKK
 3: KK KKKKKKKKKKKKKKKKK KKKKK
 4: KKK KKKKKKKKKKKKKKKKK KKKKK
 5: KKKK KKKKKKKKKKKKKKKKK KKKKK
 6: KKKKK (K) {20} KKKKK
 7: KKKKK (K) {20} KKKKK
 8: KKKKK (K) {20} KKKKK
 9: KKKKK (K) {20} KKKKK
 10: KKKKK (K) {20} KKKKK
 11: KKKKK (K) {20} KKKKK
 12: KKKKK (K) {20} KKKKK
 13: KKKKK (K) {20} KKKKK
 14: KKKKK (K) {20} KKKKK
 15: KKKKK (K) {20} KKKKK
 16: KKKKK (K) {20} KKKKK

```

17: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
18: KKKK KKKKKKKKKKKKKKKKKKK KKKK
19: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
20: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
21: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
22: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
23: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
24: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
25: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
26: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
27: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
28: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
29: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
30: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
31: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
32: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
33: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
34: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
35: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
36: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}
37: KKKK KKKKKKKKKKKKKKKKKKK KKKK
      (K) {20}

ABG7J151 ck: 8024 len: 84 | Abg7J151 Single-chain antigen-binding (sFv)
      (R,K) {20,20}
      (R) {20}
1: RRRRRRRRRRRRRRRRRR RRRR
      (R) {20}
2: R RRRRRRRRRRRRRRRR RRRR
      (R) {20}

```

[illegible]

28: RRRR (R) {20}
 29: RRRR (R) {20}
 30: RRRR (R) {20}
 31: RRRR (R) {20}
 32: RRRR (R) {20}
 33: RRRR (R) {20}
 34: RRRR (R) {20}
 35: RRRR (R) {20}
 36: RRRR (R) {20}
 37: RRRR (R) {20}

ABG73152 ck: 9643 len: 83 1 Abg73152 Single-chain antigen-binding (scFv)

1

1: RRRR (R, K) {20, 20}
 2: RRRR (R, K) {20}
 3: RRRR (R, K) {20}
 4: RRRR (R, K) {20}
 5: RRRR (R, K) {20}
 6: RRRR (R, K) {20}
 7: RRRR (R, K) {20}
 8: RRRR (R, K) {20}
 9: RRRR (R, K) {20}
 10: RRRR (R, K) {20}
 11: RRRR (R, K) {20}
 12: RRRR (R, K) {20}
 13: RRRR (R, K) {20}

14: RRRR (R, K) {20}
 15: RRRR (R, K) {20}
 16: RRRR (R, K) {20}
 17: RRRR (R, K) {20}
 18: RRRR (R, K) {20}
 19: RRRR (R, K) {20}
 20: RRRR (R, K) {20}
 21: RRRR (R, K) {20}
 22: RRRR (R, K) {20}
 23: RRRR (R, K) {20}
 24: RRRR (R, K) {20}
 25: RRRR (R, K) {20}
 26: RRRR (R, K) {20}
 27: RRRR (R, K) {20}
 28: RRRR (R, K) {20}
 29: RRRR (R, K) {20}
 30: RRRR (R, K) {20}
 31: RRRR (R, K) {20}
 32: RRRR (R, K) {20}
 33: RRRR (R, K) {20}
 34: RRRR (R, K) {20}
 35: RRRR (R, K) {20}
 36: RRRR (R, K) {20}
 37: RRRR (R, K) {20}

10: KKKKK (K) {20}
 11: KKKKK (K) {20}
 12: KKKKK (K) {20}
 13: KKKKK (K) {20}
 14: KKKKK (K) {20}
 15: KKKKK (K) {20}
 16: KKKKK (K) {20}
 17: KKKKK (K) {20}
 18: KKKKK (K) {20}
 19: KKKKK (K) {20}
 20: KKKKK (K) {20}
 21: KKKKK (K) {20}
 22: KKKKK (K) {20}
 23: KKKKK (K) {20}
 24: KKKKK (K) {20}
 25: KKKKK (K) {20}
 26: KKKKK (K) {20}
 27: KKKKK (K) {20}
 28: KKKKK (K) {20}
 29: KKKKK (K) {20}
 30: KKKKK (K) {20}
 31: KKKKK (K) {20}
 32: KKKKK (K) {20}
 33: KKKKK (K) {20}

34: KKKKK
 35: KKKKK (K) {20}
 36: KKKKK (K) {20}
 37: KKKKK (K) {20}
 ABG73871 ck: 8024 len: 84 ! Abg73871 Single-chain antigen-binding (sfv)
 1: (R, K) {20, 20}
 2: R
 3: RR
 4: RRR
 5: RRRR
 6: RRRRR
 7: RRRRR
 8: RRRRR
 9: RRRRR
 10: RRRRR
 11: RRRRR
 12: RRRRR
 13: RRRRR
 14: RRRRR
 15: RRRRR
 16: RRRRR
 17: RRRRR
 18: RRRRR
 19: RRRRR
 20: RRRRR
 21: RRRRR
 22: RRRRR
 23: RRRRR
 24: RRRRR
 25: RRRRR
 26: RRRRR
 27: RRRRR
 28: RRRRR
 29: RRRRR
 30: RRRRR
 31: RRRRR
 32: RRRRR
 33: RRRRR

295: RKKK (K) {20} KKKK
296: KGRK (K) {20} KKKK
297: GKKK (K) {20} KKKK
298: KKKK (K) {20} KKKK
299: KKKK (K) {20} KKKK
300: KKKK (K) {20} KKKK
301: KKKK (K) {20} KKKK
302: KKKK (K) {20} KKKK
303: KKKK (K) {20} KKKK
304: KKKK (K) {20} KKKK
305: KKKK (K) {20} KKKK
306: KKKK (K) {20} KKKK
342: PYKG (K) {20} KKKK
343: YNGK (K) {20} KKKK
344: KGRK (K) {20} KKKK
345: GERK (K) {20} KKKK
346: EKKK (K) {20} KKKK
347: KKKK (K) {20} KKKK
348: KKKK (K) {20} KKKK
349: KKKK (K) {20} KKKK
350: KKKK (K) {20} KKKK
351: KKKK (K) {20} KKKK
352: KKKK (K) {20} KKKK
353: KKKK (K) {20} KKKK

1

354: KKKK (K) {20} KKKK DTLT
ABJ19654 ck: 6732 len: 139 1 Abj19654 Human secreted protein amino acid
(R,K) {20,20}
(K) {20}
91: HFRPG KKKK KKKK KKKK KKKK
92: FRPG (K) {20} KKKK
93: RPKK (K) {20} KKKK
94: PKKK (K) {20} KKKK
95: GKKK (K) {20} KKKK
96: KKKK (K) {20} KKKK
97: KKKK (K) {20} KKKK
98: KKKK (K) {20} KKKK
99: KKKK (K) {20} KKKK
100: KKKK (K) {20} KKKK
101: KKKK (K) {20} KKKK
102: KKKK (K) {20} KKKK
103: KKKK (K) {20} KKKK
104: KKKK (K) {20} KKKK
105: KKKK (K) {20} KKKK
106: KKKK (K) {20} KKKK
107: KKKK (K) {20} KKKK
108: KKKK (K) {20} KKKK
109: KKKK (K) {20} KKKK
110: KKKK (K) {20} KKKK
111: KKKK (K) {20} KKKK
112: KKKK (K) {20} KKKK

113: KKKK (K) {20} KKKK
 114: KKKK (K) {20} KKKK
 115: KKKK (K) {20} KKKK
 116: KKKK (K) {20} KKKK
 117: KKKK (K) {20} KKKK
 118: KKKK (K) {20} KKKK
 119: KKKK (K) {20} K
 120: KKKK (K) {20} KKKK

ABP99475 ck: 6732 len: 139 1 Abp99475 Human secreted protein SEQ ID NO 4

1

91: HPRPG (R,K) {20,20} KKKK
 92: FRPG (K) {20} KKKK
 93: RPRG (K) {20} KKKK
 94: PRGK (K) {20} KKKK
 95: GRGK (K) {20} KKKK
 96: KKKK (K) {20} KKKK
 97: KKKK (K) {20} KKKK
 98: KKKK (K) {20} KKKK
 99: KKKK (K) {20} KKKK
 100: KKKK (K) {20} KKKK
 101: KKKK (K) {20} KKKK
 102: KKKK (K) {20} KKKK
 103: KKKK (K) {20} KKKK
 104: KKKK (K) {20} KKKK
 105: KKKK (K) {20} KKKK

106: KKKK (K) {20} KKKK
 107: KKKK (K) {20} KKKK
 108: KKKK (K) {20} KKKK
 109: KKKK (K) {20} KKKK
 110: KKKK (K) {20} KKKK
 111: KKKK (K) {20} KKKK
 112: KKKK (K) {20} KKKK
 113: KKKK (K) {20} KKKK
 114: KKKK (K) {20} KKKK
 115: KKKK (K) {20} KKKK
 116: KKKK (K) {20} KKKK
 117: KKKK (K) {20} KKKK
 118: KKKK (K) {20} KKKK
 119: KKKK (K) {20} K
 120: KKKK (K) {20} KKKK

1

ABP99639 ck: 9531 len: 66 1 Abp99639 Human secreted protein SEQ ID NO 5f
 41: MMTVX (R,K) {20,20} KKKK
 42: WTVXK (K) {20} KKKK
 43: TVXK (K) {20} KKKK
 44: VXXK (K) {20} KKKK
 45: XXXK (K) {20} KKKK
 46: KKKK (K) {20} K
 47: KKKK (K) {20} KKKK

Total finds: 8,655
Total length: 158,726,570
Total sequences: 1,107,863
CPU time: 07:06.66

```

!!SEQUENCE LIST 1.0
! FINDPATTERNS on geneseqp:* allowing 0 mismatches
!      1 C(E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V)
GENESQ2000S: AAB13781 ck: 1303 len: 25 finds: 1 ! Aab13781 Soluble peptide and
GENESQ2000S: AAB13783 ck: 4553 len: 45 finds: 1 ! Aab13783 Soluble tandem pEA/
\\End of list

Databases searched:
  Geneseq-AA, Release 13.0, Released on 19Jun2003, Formatted on 15Jul2003

Total finds:      2
Total length:    158,726,570
Total sequences: 1,107,863
CPU time:        08:18.53
  
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```

!!AA_SEQUENCE 1.0
ID AAB13781 standard; peptide; 25 AA.
XX
AC AAB13781;
XX
DT 10-NOV-2000 (first entry)
XX
DE Soluble peptide antigen pEA.
XX
KW pEA peptide; cytotoxic; vaccine; cytotoxic T cell; CTL; immunotherapy;
major histocompatibility complex class I; MHC class I; antigen; tumour;
prostate; breast; multiple myeloma.
XX
OS Unidentified.
XX
PN WO20035949-A1.
XX
PD 22-JUN-2000.
XX
PF 14-DEC-1999; 99WO-US29724.
XX
PR 14-DEC-1998; 98US-0112324.
XX
PA (DEND-) DENDREON CORP.
XX
PI Laus R, Hakim I, Vidovic D;
XX
DR WPI: 2000-442365/38.
XX
PT Antigens modified by the covalent addition of a peptide that
PT facilitates entry into antigen presenting cells, useful for producing
PT compositions for immunizing against tumors and pathogens -
XX
PS Claim 2; Page 26; 34pp; English.
XX
CC The present invention relates to compositions of modified soluble protein
CC antigens capable of eliciting an enhanced in vivo cytotoxic T cell (CTL)
CC response i.e. a major histocompatibility complex (MHC) class I molecule
CC response. The protein antigen is modified by the covalent addition of a
CC peptide sequence which facilitate entry of the antigen into antigen
CC presenting cells (APCs). The present sequence is one such peptide
CC sequence which can be used to modify the soluble antigens. The present
CC sequence is peptide pEA. The modified antigen composition may be used for
CC immunising against, or treating a tumour e.g. prostate and breast
CC carcinoma or multiple myeloma, or pathogen in mammals.
XX
SQ Sequence 25 AA;
AAB13781 Length: 25 January 30, 2004 10:59 Type: P Check: 1303 ..
1 CEAATAAATAA AAAAATAAAE AAAAA
!!AA_SEQUENCE 1.0
ID AAB13783 standard; peptide; 45 AA.
XX
AC AAB13783;
XX
DT 10-NOV-2000 (first entry)
XX
DE Soluble tandem pEA/ pK peptide conjugate.
XX
KW pK peptide; cytotoxic; vaccine; cytotoxic T cell; CTL; immunotherapy;
major histocompatibility complex class I; MHC class I; antigen; tumour;
prostate; breast; multiple myeloma; pEA peptide.
XX
OS Unidentified.
XX
PN WO20035949-A1.
XX
PD 22-JUN-2000.
XX
PF 14-DEC-1999; 99WO-US29724.
XX

```

```

PR 14-DEC-1998; 98US-0112324.
XX
PA (DEND-) DENDREON CORP.
XX
PI Laus R, Hakim I, Vidovic D;
XX
DR WPI: 2000-442365/38.
XX
PT Antigens modified by the covalent addition of a peptide that
PT facilitates entry into antigen presenting cells, useful for producing
PT compositions for immunizing against tumors and pathogens -
XX
PS Claim 2; Page 26; 34pp; English.
XX
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CC antigens capable of eliciting an enhanced in vivo cytotoxic T cell (CTL)
CC response i.e. a major histocompatibility complex (MHC) class I molecule
CC response. The protein antigen is modified by the covalent addition of a
CC peptide sequence which facilitate entry of the antigen into antigen
CC presenting cells (APCs). The present sequence is one such peptide
CC sequence which can be used to modify the soluble antigens. The present
CC sequence is tandem pEA/ pK peptide conjugate. The modified antigen
CC composition may be used for immunising against, or treating a tumour e.g.
CC prostate and breast carcinoma or multiple myeloma, or pathogen in
CC mammals.
XX
SQ Sequence 45 AA;
AAB13783 Length: 45 January 30, 2004 10:59 Type: P Check: 4553 ..
1 CEAATAAATAA AAAAATAAAE AAAAAKKKK KKKKKKKKK KKKKK

```


! FINDPATTERNS on pir:* allowing 0 mismatches

! 1 C(E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V)

Databases searched:

NBRF, Release 76.1, Released on 12May2003, Formatted on 10Jun2003

Total finds:

0

Total length:

96,168,682

Total sequences:

283,308

CPU time:

02:26.82

! FINDPATTERNS on swp:* allowing 0 mismatches

! 1 C(E,D) (A,L,I,F,G,C,M,V){5,5} (B,D) (A,L,I,F,G,C,M,V){5,5} (B,D) (A,L,I,F,G,C,M,V)

Databases searched:

SWISS-PROT, Release 41.1, Released on 6Jun2003, Formatted on 9Jun2003
 SPTREMBL, Release 23.0, Released on 4Mar2003, Formatted on 7Mar2003

Total finds: 0
 Total length: 305,079,309
 Total sequences: 958,388
 CPU time: 08:14.71

! FINDPATTERNS on geneseqp: * allowing 0 mismatches

1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

AAB13781 ck: 1303 len: 25 ! Aab13781 Soluble peptide antigen p5A. 11/20

1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

2: C EAAAAAAAAAAAAAAAAA EAAAA

8: AAAAA EAAAAAAAAAAAAAAAAA

AAB13783 ck: 4553 len: 45 ! Aab13783 Soluble tandem p5A/ PK peptide con

1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

2: C EAAAAAAAAAAAAAAAAA EAAAA

8: AAAAA EAAAAAAAAAAAAAAAAA KKKKK

ABP02760 ck: 5947 len: 86 ! Abp02760 Human ORFX protein sequence SEQ ID

1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

13: GGRVR DACGCCEVCGALCGAVCG LQEGP

Databases searched:
Geneseq-AA, Release 13.0, Released on 19Jun2003, Formatted on 15Jul2003

Total finds: 5
Total length: 158,726,570
Total sequences: 1,107,863
CPU time: 09:16.70


```

!!SEQUENCE LIST 1.0
! FINDPATTERNS on geneseqp:* allowing 0 mismatches
!
! 1 (E,D) (A,L,I,F,G,C,M,V){5,5} (E,D) (A,L,I,F,G,C,M,V) {
GENESEQP2000S:AAB13781 ck: 1303 len: 25 finds: 2 ! Aab13781 Soluble peptide ant
GENESEQP2000S:AAB13783 ck: 4553 len: 45 finds: 2 ! Aab13783 Soluble tandem pEA/
GENESEQP2002S:ABP02760 ck: 5947 len: 86 finds: 1 ! ABP02760 Human ORFX protein
\\End of list

Databases searched:
Geneseq-AA, Release 13.0, Released on 19Jun2003, Formatted on 15Jul2003

Total finds: 5
Total length: 158,726,570
Total sequences: 1,107,863
CPU time: 11:32.04

```


!!AA_SEQUENCE 1.0
ID AAB13781 standard; peptide; 25 AA.
XX
AC AAB13781;
XX
DT 10-NOV-2000 (first entry)
XX
DE Soluble peptide antigen pEA.
XX
KM pEA peptide; cytotoxic; vaccine; cytotoxic T cell; CTL; immunotherapy;
KM major histocompatibility complex class 1; MHC class 1; antigen; tumour;
KM prostate; breast; multiple myeloma.
XX
OS Unidentified.
XX
PN WO200035949-A1.
XX
PD 22-JUN-2000.
XX
PF 14-DEC-1999; 99WO-US29724.
XX
PR 14-DEC-1998; 98US-0112324.
XX
PA (DEND-) DENDREON CORP.
XX
PI Laus R, Hakim I, Vidovic D;
XX
DR WPI; 2000-442365/38.
XX
PT Antigens modified by the covalent addition of a peptide that
PT facilitates entry into antigen presenting cells, useful for producing
PT compositions for immunizing against tumors and pathogens -
XX
PS Claim 2; Page 26; 34pp; English.
XX
CC The present invention relates to compositions of modified soluble protein
CC antigens capable of eliciting an enhanced in vivo cytotoxic T cell (CTL)
CC response i.e. a major histocompatibility complex (MHC) class I molecule
CC response. The protein antigen is modified by the covalent addition of a
CC peptide sequence which facilitate entry of the antigen into antigen
CC presenting cells (APCs). The present sequence is one such peptide
CC sequence which can be used to modify the soluble antigens. The present
CC sequence is peptide pEA. The modified antigen composition may be used for
CC immunizing against, or treating a tumour e.g. prostate and breast
CC carcinoma or multiple myeloma, or pathogen in mammals.
XX
SQ Sequence 25 AA;
XX
AAB13781 Length: 25 January 30, 2004 11:00 Type: P Check: 1303 ..
1 CEAATAAATAA AAAATAAAE AAAA
!!AA_SEQUENCE 1.0
ID AAB13783 standard; peptide; 45 AA.
XX
AC AAB13783;
XX
DT 10-NOV-2000 (first entry)
XX
DE Soluble tandem pEA/ pK peptide conjugate.
XX
KM pK peptide; cytotoxic; vaccine; cytotoxic T cell; CTL; immunotherapy;
KM major histocompatibility complex class 1; MHC class 1; antigen; tumour;
KM prostate; breast; multiple myeloma; pEA peptide.
XX
OS Unidentified.
XX
PN WO200035949-A1.
XX
PD 22-JUN-2000.
XX
PF 14-DEC-1999; 99WO-US29724.
XX

PR 14-DEC-1998; 98US-0112324.
XX
PA (DEND-) DENDREON CORP.
XX
PI Laus R, Hakim I, Vidovic D;
XX
DR WPI; 2000-442365/38.
XX
PT Antigens modified by the covalent addition of a peptide that
PT facilitates entry into antigen presenting cells, useful for producing
PT compositions for immunizing against tumors and pathogens -
XX
PS Claim 2; Page 26; 34pp; English.
XX
CC The present invention relates to compositions of modified soluble protein
CC antigens capable of eliciting an enhanced in vivo cytotoxic T cell (CTL)
CC response i.e. a major histocompatibility complex (MHC) class I molecule
CC response. The protein antigen is modified by the covalent addition of a
CC peptide sequence which facilitate entry of the antigen into antigen
CC presenting cells (APCs). The present sequence is one such peptide
CC sequence which can be used to modify the soluble antigens. The present
CC sequence is tandem pEA/ pK peptide conjugate. The modified antigen
CC composition may be used for immunising against, or treating a tumour e.g.
CC prostate and breast carcinoma or multiple myeloma, or pathogen in
CC mammals.
XX
SQ Sequence 45 AA;
XX
AAB13783 Length: 45 January 30, 2004 11:00 Type: P Check: 4553 ..
1 CEAATAAATAA AAAATAAAE AAAAAXKKK KKKKKKKK KKKK
!!AA_SEQUENCE 1.0
ID AAB02760 standard; Protein; 86 AA.
XX
AC AAB02760;
XX
DT 25-JUN-2002 (first entry)
XX
DE Human ORFX protein sequence SEQ ID NO:5502.
XX
XX Human; open reading frame; ORFX; gene therapy; cancer; cirrhosis;
XX hyperproliferative disorder; psoriasis; benign tumour; haemorrhage;
XX degenerative disorder; osteoarthritis; neurodegenerative disorder;
XX cardiovascular disease; diabetes mellitus; systemic lupus erythematosus;
XX hypertension; hypothyroidism; cholesterol ester storage disease;
XX immune deficiency; immune disorder; infectious disease;
XX autoimmune disorder; rheumatoid arthritis; autoimmune thyroiditis;
XX myaethenia gravis.
XX
OS Homo sapiens.
XX
PN WO200192523-A2.
XX
PD 06-DEC-2001.
XX
PF 29-MAY-2001; 2001WO-US10836.
XX
PR 30-MAY-2000; 2000US-206132P.
XX
PR 29-AUG-2000; 2000US-228716P.
XX
PA (CURA-) CURAGEN CORP.
XX
PI Shinkets RA, Leach MD;
XX
DR N-PSDB; AEN18512.
XX
WPI; 2002-106308/14.
XX
N-PSDB; AEN18512.
XX
PT Novel human polypeptides and polynucleotides useful for diagnosing,
PT preventing and treating cardiovascular disease, neurodegenerative,
PT hyperproliferative disorders and autoimmune disorders -
XX
PS Disclosure; SEQ ID 5502; 1037pp; English.

XX The present invention describes substantially purified human proteins
 CC (referred to as open reading frame, ORFX, where X is 1-11491 (see Table 1
 CC in the specification). ABN15762 to ABN27252 encode the human ORFX
 CC proteins given in ABP00010 to ABP11500. ORFX proteins are useful for
 CC treating or preventing a pathology associated with an ORFX-associated
 CC disorder in humans, and in the manufacture of a medicament for treating a
 CC syndrome associated with ORFX-associated disorder. ORFX polynucleotide
 CC sequences can be used in gene therapy. ORFX sequences can be used in the
 CC treatment of cancer, hyperproliferative disorders, cirrhosis of liver,
 CC psoriasis, benign tumours, keloid, degenerative disorders, haemorrhage,
 CC osteoarthritis, neurodegenerative disorders, disorders related to organ
 CC transplantation, cardiovascular diseases, diabetes mellitus, systemic
 CC lupus erythematosus, hypertension, hypothyroidism, cholesterol ester
 CC storage disease, various immune deficiencies and disorders, infectious
 CC diseases, autoimmune disorders such as multiple sclerosis, rheumatoid
 CC arthritis, autoimmune thyroiditis, myasthenia gravis, graft-versus-host
 CC disease and autoimmune inflammatory eye disease. ORFX proteins are also
 CC useful for treating burns, incisions, ulcers, for treating osteoporosis,
 CC bone degenerative disorders, or periodontal disease, and for gut
 CC protection or regeneration and treatment of lung or liver fibrosis,
 CC reperfusion injury in various tissues and conditions resulting from
 CC systemic cytokine damage.
 CC N.B. The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 CC

XX
 SQ Sequence 86 AA;

ABP02760 Length: 86 January 30, 2004 11:00 Type: P Check: 5947 ..

1 XPPTDCEGR VADACGCEV CGALEGAVCG LQEGPCGEA ANAVSAPSG

51 VPASATVRK AQAGLCVCAS SEPVGNDK TYTNLC

! FINDPATTERNS on swp:* allowing 0 mismatches

1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

HRAL_MOUSE ck: 7206 len: 480 ! Q9r118 mus musculus (mouse). serine protease

1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

60: GGRVR DACGCCVCGALGGAACG LQEGP

1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

Q9KHD9 ck: 1697 len: 262 ! Q9khd9 streptomyces griseus subsp. griseus.
(E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

124: LGGGL ELALACDLVAVAGSALFG LPFLG

1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

Q8RUS6 ck: 3612 len: 107 ! Q8rus6 oryza sativa (japonica cultivar-group)
(E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

20: LETEV EAIAIVEGGCGVDLAVG RALGL

1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

Q9QZK6 ck: 7594 len: 480 ! Q9qzk6 mus musculus (mouse). insulin-like growth factor 1
(E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

60: GGRVR DACGCCVCGALGGAACG LQEGP

1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

Q91WS3 ck: 7882 len: 480 ! Q91ws3 mus musculus (mouse). protease, serine
(E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

60: GGRVR DACGCCVCGALGGAACG LQEGP

1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

Q9QZK5 ck: 8689 len: 480 ! Q9qzk5 rattus norvegicus (rat). insulin-like growth factor 1
(E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {5,5}

60: GGRVR DACGCCVCGALGGAACG LQEGP

Databases searched:

SWISS-PROT, Release 41.1, Released on 6Jun2003, Formatted on 9Jun2003
SPTREMBL, Release 23.0, Released on 4Mar2003, Formatted on 7Mar2003

Total finds: 6

Total length: 305,079,309

Total sequences: 958,388

CPU time: 14:58.50

!!SEQUENCE LIST 1.0
! FINDPATTERNS on swp: * allowing 0 mismatches

! 1 (E,D) (A,L,I,F,G,C,M,V) {5,5} (E,D) (A,L,I,F,G,C,M,V) {

SW:HRAL_MOUSE	ck: 7206	len: 480	finds: 1	! Q9r118	mus musculus (mouse) . S
SP_BA:Q9KHD9	ck: 1697	len: 262	finds: 1	! Q9khd9	streptomyces griseus su
SP_PU:Q8RUS6	ck: 3612	len: 107	finds: 1	! Q8rus6	oryza sativa (japonica
SP_RO:Q9QZK6	ck: 7594	len: 480	finds: 1	! Q9qzk6	mus musculus (mouse) . I
SP_RO:Q91WS3	ck: 7882	len: 480	finds: 1	! Q91ws3	mus musculus (mouse) . F
SP_RO:Q9QZK5	ck: 8689	len: 480	finds: 1	! Q9qzk5	rattus norvegicus (rat)

\\End of list

Databases searched:

SWISS-PROT, Release 41.1, Released on 6Jun2003, Formatted on 9Jun2003
SPRTRMBL, Release 23.0, Released on 4Mar2003, Formatted on 7Mar2003

Total finds: 6
Total length: 305,079,309
Total sequences: 958,388
CPU time: 16:49.45


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!!AA_SEQUENCE 1.0
ID HRAI_MOUSE STANDARD; PRT; 480 AA.
AC OPR18;
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Serine protease HTRA1 precursor (EC 3.4.21.-).
GN PRSS11 OR HTRA1 OR HTRA1.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ICR; TISSUE=Brain;
RA Oka C., Soma A., Kanda H., Kawachi M.;
RT "The role of murine serine protease HTRA in osteogenesis.";
RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: Protease that regulate the availability of IGFs by
CC cleaving IGF-binding proteins (By similarity).
CC -1- SUBCELLULAR LOCATION: Secreted (By similarity).
CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY S2C.
CC -1- SIMILARITY: Contains 1 IGFBP domain.
CC -1- SIMILARITY: Contains 1 Kazal-like domain.
CC -1- SIMILARITY: Contains 1 PDZ/DHR domain.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.ebi.ac.uk/announcements/
CC or send an email to license@ebi.ac.uk).
CC -----
DR EMBL; AF172994; AAD9422.1; -.
DR MEROPS; S01.277; -.
DR MGD; MGI:1929076; Prss11.
DR InterPro; IPR000867; Insl_gro_fac_pr.
DR InterPro; IPR002350; Kazal.
DR InterPro; IPR001478; PDZ.
DR InterPro; IPR001940; Protease2C.
DR InterPro; IPR001254; Ser_protease_Try.
DR Pfam; PF00050; Kazal; 1.
DR Pfam; PF00595; PDZ; 1.
DR Pfam; PF00089; trypsin; 1.
DR PRINTS; PR00834; PROTEASE2C.
DR SMART; SM00121; IB; 1.
DR SMART; SM00280; KAZAL; 1.
DR SMART; SM00228; PDZ; 1.
DR PROSITE; PS50106; PDZ; 1.
KW Hydrolyase; Serine protease; Growth factor binding; Signal.
FT SIGNAL 1 22 POTENTIAL.
FT CHAIN 1 480 SERINE PROTEASE HTRA1.
FT DOMAIN 23 37 IGFBP.
FT DOMAIN 37 94 KAZAL-LIKE.
FT DOMAIN 101 155 SERINE PROTEASE.
FT DOMAIN 204 364 PDZ.
FT ACT_SITE 365 467 CHARGE RELAY SYSTEM (POTENTIAL).
FT ACT_SITE 220 220 CHARGE RELAY SYSTEM (POTENTIAL).
FT ACT_SITE 250 250 CHARGE RELAY SYSTEM (POTENTIAL).
FT ACT_SITE 328 328 CHARGE RELAY SYSTEM (POTENTIAL).
SQ SEQUENCE 480 AA; 51246 MW; 54BB9BA6C99A7BF4 CRC64;
HRAI_MOUSE Length: 480 January 30, 2004 15:08 Type: P Check: 7206
1 M0SLRTTLLS LLLILLAPS LALPSTGTRS APAATVCPEN CDPTRCAPP
51 TDCBGRVND ACGCEVCGA LEGACGLQE GPCGEGLCV LPFGVPASAT
101 VRRRAQAGC VCASPEVCG SDAKTTNLC QLRASRRE KLPPQPVIVL
151 ORGACGQOE DPNLRHKYN FIADVKEFA PDVVKHLYR KLPPSKREV
201 VASGGFIVS EDGLIVTNAH VTNKRVKV ELKNGATYEA IIKVDEKAD

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!!AA_SEQUENCE 1.0
ID Q9KHD9 PRELIMINARY; PRT; 262 AA.
AC Q9KHD9;
DT 01-OCT-2000 (Tremblrel. 15, Created)
DT 01-OCT-2000 (Tremblrel. 15, Last sequence update)
DT 01-MAR-2002 (Tremblrel. 20, Last annotation update)
DE Enoyl-CoA hydratase-like protein.
OS Streptomyces griseus subsp. griseus.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Streptomycinae; Streptomycetaceae; Streptomyces.
OX NCBI_TaxID=67263;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=DSM40695;
RX MEDLINE=20316791; PubMed=10858335;
RA Smith W.C., Xiang L., Shen B.;
RT "Genetic localization and molecular characterization of the nons gene
RT required for macroreticoid biosynthesis in Streptomyces griseus
RT DSM40695.";
RL Actinmicrob. Agents Chemother. 44:1809-1817(2000).
DR EMBL; AF263011; AAF81231.1; -.
DR HSSP; P14604; 2DUB.
DR InterPro; IPR001753; EnCoA_hydrase.
DR Pfam; PF00378; ECH; 1.
DR PROSITE; PS00166; ENOYL_COA_HYDRATASE; 1.
SQ SEQUENCE 262 AA; 27068 MW; D218PA2BDEG64BF2 CRC64;
Q9KHD9 Length: 262 January 30, 2004 15:08 Type: P Check: 1697
1 MTPETAPBEG EBPPLAVERR GRVATLTLLNR PHRRNANSTA MLARLDHALG
51 KLTLQGGGAP GALVTGAGG TFSGGADTRE PDWDLSSRA VRRAHPRTVF
101 AMLEHAPPPV VAAVEGYALG GGLIELALCD LVVAGEGALF GLPELVGAV
151 PGGAHVSLV RRAGRGVNAR MLITGERYR ADELARLGA VERTVDDGAL
201 AEAQALASV AAGDPALLBA GVRLRDSGH LDRTAALGVE NGYWMQASA
251 ANRDPSPSSG RP
!!AA_SEQUENCE 1.0
ID Q8RUS6 PRELIMINARY; PRT; 107 AA.
AC Q8RUS6;
DT 01-JUN-2002 (Tremblrel. 21, Created)
DT 01-JUN-2002 (Tremblrel. 21, Last sequence update)
DT 01-OCT-2002 (Tremblrel. 22, Last annotation update)
DE P0678F11.3 protein (P0413C03.30 protein).
GN P0678F11.3 OR P0413C03.30.
OS Oryza sativa (japonica cultivar-group).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Eriocaridaceae; Oryzaceae; Oryza.
OX NCBI_TaxID=39947;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Nipponbare;
RA Sasaki T., Matsumoto T., Yamamoto K.;
RT "Oryza sativa nipponbare (GA3) genomic DNA, chromosome 1, PAC
RL clone:P0678F11.";
RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.
[2]

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RP SEQUENCE FROM N.A.
RC STRAIN=cv. Nipponbare;
RA Sasaki T., Matsumoto T., Yamamoto K.;
RT "Oryza sativa nipponbare (G3) genomic DNA, chromosome 1, PAC
   clone:p0413C03."
RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL; AP003437; BAB86096.1; -
DR EMBL; AP003451; BAB86154.1; -
DR Gramene; Q8RUS6; -
SQ SEQUENCE 107 AA; 11046 MW; 89097D88201BD849 CRC64;

Q8RUS6 Length: 107 January 30, 2004 15:08 Type: P Check: 3612 ..

1 MESPRAVNL RMSLETEVE AALAVEGCG VLLAWGRAL GLDPATVALN
51 GYFVRGKGH VSAVTWRAL LDFPARGLP TGDAPAPVA VHGKAPPPP
101 PPPVSVL

!!AA SEQUENCE 1.0
ID _Q9QZK6 PRELIMINARY; PRT; 480 AA.
AC Q9QZK6;
DT 01-MAY-2000 (TREMBlrel. 13, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE Insulin-like growth factor binding protein 5 protease.
GN PRS11.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6; TISSUE=Ovary;
RA Houtvitz A., Hennebold J.D., King G., Negishi H., Erickson G.F.,
RA Rody J.A., Mayo K.E., Adashi E.Y.;
RT "Mouse insulin-like growth factor binding protein 5-directed
   endopeptidase: structural assessment, evolutionary analysis, ovarian
   expression, hormonal regulation and cellular localization.";
RL Submitted (AUG-1999) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: CONTAINS 1 PDZ/DHR DOMAIN.
DR EMBL; AF179369; AAD52682.1; -
DR MGD; MGI:1929076; Prs11.
DR InterPro; IPR000867; Insl_gro_fac_pr.
DR InterPro; IPR002350; kazal.
DR InterPro; IPR001478; PDZ.
DR InterPro; IPR001940; Protease2C.
DR InterPro; IPR001254; Ser_protease_Try.
DR Pfam; PF00219; IGFBP_1.
DR Pfam; PF00595; kazal_1.
DR Pfam; PF00595; PDZ_1.
DR Pfam; PF00089; trypsin_1.
DR PRINTS; PRO0834; PROTEASES2C.
DR SMART; SM00121; IB; 1.
DR SMART; SM00280; KAZAL; 1.
DR SMART; SM00228; PDZ_1.
DR SMART; PS50106; PDZ_1.
DR PROSITE; PS50106; PDZ; 1.
DR Hydrilase; Protease; Serine protease.
KW SEQUENCE 480 AA; 51213 MW; 92BDDA85CF5B1287 CRC64;

Q9QZK6 Length: 480 January 30, 2004 15:08 Type: P Check: 7594 ..

1 MSLRTTLLS LLLLLLAPS LALPSGTGRS APAATVCPFH CDPTRCAPPP
51 TDCEGRVRD AGCCCEVCGA LEGAAGLQGE GPCGEGLOCV VPGVPASAT
101 VRRRAQAGLC VCASSEPVCG SDAKTYTNLC QLRASRSE KLRPPVIVL
151 QRGACGQGE DPNLSRHKN FIAVVEKIA PAVVHIELYR KLPSKREVP
201 VASGSGFIVS EDGLIVTNAH VTNKNRKVY ELKNGATVEA KIKDVEDKAD
251 IALIKIDHOG KLPVLLGRS SELRPGFVV AIGSPFSLQN TVTTGIVSTT

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301 QRGKEIGLAR NSDMYIQTQ AIINYNSGG PLVNLDEVI GINTLKVTAG
351 ISFALPSFKI KKFLETSHDR QAKGKAVTK KYIGRMSL TSSAKELKD
401 RHRDPDVLS GAVIIEVIPD TPAEAGLKE NDVIISINGQ SVTANDVSD
451 VIKKENTILM VVRGNEDIV ITVPEIDP

!!AA SEQUENCE 1.0
ID _Q91WS3 PRELIMINARY; PRT; 480 AA.
AC Q91WS3;
DT 01-DEC-2001 (TREMBlrel. 19, Created)
DT 01-DEC-2001 (TREMBlrel. 19, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE Protease, serine, 11 (1gf binding).
GN PRS11.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Kidney;
RA Strauberg R.;
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: CONTAINS 1 PDZ/DHR DOMAIN.
DR EMBL; BC013516; AAH13516.1; -
DR MGD; MGI:1929076; Prs11.
DR InterPro; IPR000867; Insl_gro_fac_pr.
DR InterPro; IPR002350; kazal.
DR InterPro; IPR001478; PDZ.
DR InterPro; IPR001940; Protease2C.
DR InterPro; IPR001254; Ser_protease_Try.
DR Pfam; PF00219; IGFBP_1.
DR Pfam; PF00595; kazal_1.
DR Pfam; PF00595; PDZ_1.
DR PRINTS; PRO0834; PROTEASES2C.
DR SMART; SM00121; IB; 1.
DR SMART; SM00280; KAZAL; 1.
DR SMART; SM00228; PDZ_1.
DR PROSITE; PS50106; PDZ; 1.
DR Hydrilase; Protease; Serine protease.
KW SEQUENCE 480 AA; 51212 MW; 76BDD5B862EDC9DA CRC64;

Q91WS3 Length: 480 January 30, 2004 15:08 Type: P Check: 7882 ..

1 MSLRTTLLS LLLLLLAPS LALPSGTGRS APAATVCPFH CDPTRCAPPP
51 TDCEGRVRD AGCCCEVCGA LEGAAGLQGE GPCGEGLOCV VPGVPASAT
101 VRRRAQAGLC VCASSEPVCG SDAKTYTNLC QLRASRSE KLRPPVIVL
151 QRGACGQGE DPNLSRHKN FIAVVEKIA PAVVHIELYR KLPSKREVP
201 VASGSGFIVS EDGLIVTNAH VTNKNRKVY ELKNGATVEA KIKDVEDKAD
251 IALIKIDHOG KLPVLLGRS SELRPGFVV AIGSPFSLQN TVTTGIVSTT
301 QRGKEIGLAR NSDMYIQTQ AIINYNSGG PLVNLDEVI GINTLKVTAG
351 ISFALPSFKI KKFLETSHDR QAKGKAVTK KYIGRMSL TSSAKELKD
401 RHRDPDVLS GAVIIEVIPD TPAEAGLKE NDVIISINGQ SVTANDVSD
451 VIKKENTILM VVRGNEDIV ITVPEIDP

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!!AA SEQUENCE 1.0
ID _Q9QZK5 PRELIMINARY; PRT; 480 AA.
AC Q9QZK5;
DT 01-MAY-2000 (TREMBlrel. 13, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)

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DT 01-MAR-2003 (TrEMBLrel. 23, last annotation update)
 DE Insulin-like growth factor binding protein 5 protease.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OX NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=Sprague-Dawley; TISSUE=Ovary;
 RA Hourvitz A., Hennebold J.D., King G., Negishi H., Erickson G.F.,
 RA Roby J.A., Mayo K.E., Adashi E.Y.;
 RT "Mouse insulin-like growth factor binding protein 5-directed
 RT endopeptidase: structural assessment, evolutionary analysis, ovarian
 RT expression, hormonal regulation and cellular localization";
 RL Submitted (Aug-1999) to the EMBL/GenBank/DBJ databases.
 CC -1- SIMILARITY: CONTAINS 1 PDZ/DHR DOMAIN.
 DR EMBL: AF179370; AAD52683.1; -.
 DR MEROPS; S01.277; -.
 DR InterPro: IPR000867; Ins1_gro_fac_pr.
 DR InterPro: IPR002350; Kazal.
 DR InterPro: IPR001478; PDZ.
 DR InterPro: IPR001940; Protease2C.
 DR InterPro: IPR01254; Ser_protease_Try.
 DR Pfam; PF00219; IGFBP; 1.
 DR Pfam; PF00050; Kazal; 1.
 DR Pfam; PF00595; PDZ; 1.
 DR Pfam; PF00089; trypsin; 1.
 DR PRINTS; PR00834; PROTEASES2C.
 DR SMART; SM00121; IB; 1.
 DR SMART; SM00280; KAZAL; 1.
 DR SMART; SM00228; PDZ; 1.
 DR PROSITE; PS50106; PDZ; 1.
 KM Hydrolyase; Protease; Serine protease.
 SQ SEQUENCE 480 AA; 5130 MW; 37A864C5A8FC035 CRC64;

Q9GZK5 Length: 480 January 30, 2004 15:08 Type: P Check: 8689 ..
 1 MQFLRTALLS LLLLLLAAPS LALPSGISRS APAATVCPFH CDPTRCAPP
 51 TDCBGRGRYRD ACGCEVCGA LBGAVCGLQE GPGCEGLQCV VPFQVPASAT
 101 VRRRAQAGLC VCASSEPCVG SDAKTYTNLC QLRPAARRSE KLRQPPVIVL
 151 QRGACGQGOE DPNGLRHKYN FIADVVEKIA PAVVHIELYR KLPSKREVP
 201 VASGSGFIVS EDGLIVTNAH VTNKRNRYK ELKNGATIEA KIKVDEKAD
 251 IALIKIDHOG KLPVLLGERS SELRPGFVV AIGSPPSLQN TVTTGIVSTT
 301 QRGKELGLR NSDMDIQTD AINYNSSG PLVNLGDEVI GINTLKVTAG
 351 ISFALPSDKI KKLFTESHDR QAKGTVTKK KYIGIRMSL TSSKAKELKD
 401 RHRDEPDIYS GAYLIEVIPT TPAAGGLKE NDVIISINQ SVVTANDVSD
 451 VIKKENTLNM VVRNGNEDIV ITVPEEIDP

! FINDPATTERNS on pir:* allowing 0 mismatches

! 1 (E,D) (A,L,I,F,G,C,M,V){5,5} (E,D) (A,L,I,F,G,C,M,V) {

Databases searched:

NBRF, Release 76.1, Released on 12May2003, Formatted on 10Jun2003

Total finds:

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Total length:

96,168,682

Total sequences:

283,308

CPU time:

04:36.11

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